

# Instructions For Use

# A00045-IFU-RUO

Revision: 5

P.O. Box 3286 - Logan, Utah 84323, U.S.A. - Tel. (800) 729-8350 - Tel. (435) 755-9848 - Fax (435) 755-0015 - www.ScyTek.com

# Vimentin; Clone V9 (Ready-To-Use)

 Catalog Number
 Volume

 A00045-0002
 2 ml

 A00045-0007
 7 ml

 A00045-0025
 25 ml

**Description** 

Species: Mouse

Immunogen: Porcine eye lens.

Clone: V9

**Isotype:** IgG1, Kappa.

Format: This antibody has been pretitered and quality controlled to work on formalin-fixed paraffin-embedded as well as acetone fixed

on formalin-fixed paramin-embedded as well as acetone fix cryostat tissue sections. No further titration is required.

Specificity: This antibody reacts with a 58kDa protein identified as vimentin.

It shows no cross-reactivity with other closely related intermediate

It shows no cross-reactivity with other closely related intermediate filament proteins such as desmin, keratin, neurofilament, and glial

fibrillary acid protein.

Species Reactivity: Reacts with Human, Rat, Horse, Chicken, Cow, Cat, Dog, and

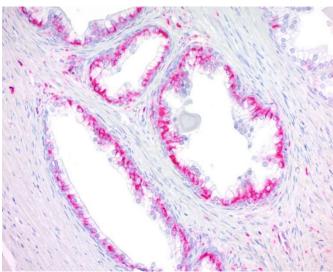
Pig. Does not react with Mouse.

Positive Control: U-87, Raji, Jurkat or HeLa cells. Lymph node or tonsil.

Cellular Localization: Cytoplasmic

Titer/Working Dilution: Ready-to-Use (no further dilution required)

Microbiological State: Nonsterile.



Human prostate stained using Vimentin; Clone V9. Pretreatment with EDTA - Saline Buffer (10X Concentrate); pH 8.0 for 5 minutes at >95°C followed by cooling to room temperature for 20 minutes. Results were visualized using ScyTek's PolyTek Anti-Mouse Polymerized Alk-Phos (Permanent Red) Staining System. Magnification 200X.

# Intended Use

Rev. Date: Feb. 12, 2021

For Research Use Only. This antibody is intended for the qualitative visualization of the anatomical elements listed in the Specificity section. It is intended to be used within an Immunohistochemistry (IHC) procedure on formalin-fixed paraffin-embedded (FFPE) human tissue followed by visualization by light microscopy. Any diagnostic interpretation of the results of this antibody is to 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.

Page 1 of 2

#### **Procedure**

- 1. Tissue Section Pretreatment (Recommended): Staining of formalin fixed, paraffin embedded tissue sections is significantly enhanced by pretreatment with EDTA Saline Buffer (10X Concentrate); pH 8.0 (ScyTek catalog# ETA500) for 5-10 minutes at >95°C followed by cooling to room temperature for 20 minutes.
- 2. **Primary Antibody Incubation Time:** We suggest an incubation period of 30 minutes at room temperature. However, depending upon the fixation conditions and the staining system employed, optimal incubation should be determined by the user.
- 3. **Visualization:** For maximum staining intensity we recommend the "CRF Anti-Polyvalent HRP Polymer" (ScyTek catalog# ABZ125, see IFU for instructions) combined with the "DAB Chromogen/Substrate Bulk Pack (High Contrast)" (ScyTek catalog# ACV500, see IFU for instructions).

# Materials and Reagents Required but not Provided

- 1. Control tissue and reagents
- 2. Xylene, graded alcohols, and deionized/distilled water
- 3. IHC detection system. Suggested: ScyTek Cat# ABZ125 "CRF Anti-Polyvalent HRP Polymer" and ScyTek Cat# ACV500 "DAB Chromogen/Substrate Kit (High Contrast)".
- 4. Wash buffer for rinses (ScyTek Cat# TBT500)
- 5. Retrieval solution (ScvTek Cat# ETA500)
- 6. Hematoxylin counterstain and bluing reagent (ScyTek Cat# HMM500 and BRT500)
- 7. Mounting medium and coverslips

**Note:** ScyTek Laboratories has a wide range of IHC reagents and ancillaries that can be found at scytek.com.

# **Storage and Stability**

Do not Freeze. Store at 2-8°C. Return to 2-8° immediately after use. Do not use after expiration date printed on label. Verify visually that antibody has not been contaminated before use. Do not use if reagent becomes cloudy or precipitates.

## Limitations

Immunohistochemistry is a complex technique involving both histological and immunological detection methods. Tissue processing and handling prior to immunostaining can cause inconsistent results. Variations in fixation and embedding or the inherent nature of the tissue specimen may cause variations in results. Endogenous peroxidase activity or pseudoperoxidase activity in erythrocytes and endogenous biotin may cause non-specific staining depending on detection system used. This data sheet's recommendations and procedures were validated using ScyTek IHC reagents and may not be suitable for other detection systems.

## **Precautions**

1. Contains Sodium Azide as a preservative (0.09% w/v), do not ingest. 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. This product contains no hazardous material at a reportable concentration according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC. 2. Do not pipette by mouth.



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Rev. Date: Feb. 12, 2021 **Revision: 5** Page 2 of 2

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- 3. Avoid contact of reagents and specimens with skin and mucous membranes.
- 4. Avoid microbial contamination of reagents or increased nonspecific staining may occur.
- The user must validate any procedures and recommendations that differ from this data sheet.
- 6. The SDS may be found at scytek.com

#### References

- Yazgan B, Sozen E, Karademir B, Ustunsoy S, Ince U, Zarkovic N, Ozer NK. CD36 expression in peripheral blood mononuclear cells reflects the onset of atherosclerosis. BioFactors. 2018 Nov;44(6):588-96.
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- Sfacteria A, Macrì F, Perillo L, Rapisarda G, Lanteri G, Mazzullo G. Cytologic and histologic features of spinal cord ependymoma in a young dog: a case report. Veterinární medicína. 2010 Feb 18;55(1):35-8.

#### <u> Warranty</u>

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