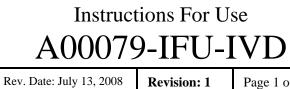
ScyTek Laboratories



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

Laminin

Availability/Contents:		<u>Item #</u> A20079 A00079 A00079.25	<u>Volume</u> 2 ml 6 ml 25 ml
Description:			
Species: Immunogen: Clone: Isotype: Format: Specificity: Species Reactivity: Positive Control: Cellular Localization: Titer/Working Dilution: Microbiological State:		Mouse A chain of human laminin. 4C7 IgG _{2a} , kappa This antibody has been pretitered and quality controlled to work on formalin-fixed paraffin- embedded as well as acetone fixed cryostat tissue sections. No further titration is required. Laminin stains the basement membrane strongly, with some weaker staining of the cytoplasm of epithelia, endothelia and smooth muscle. Human. Not reactive with mouse or rat. Intact basement membrane, Skin, Lung, Kidney or Tonsil Basement Membrane No further dilution is required. This product is not sterile.	
Uses/Limitations: For In Vitro Diagnostic Use. This product is intended for qualitative immunohistochemistry with normal and neoplastic formalin- fixed, paraffin-embedded tissue sections, to be viewed by light microscopy. Do not use past expiration date.			
Storage and Stability: 2-8° Centigrade. Product is stable for 24 months from d If reagent is not stored as recommend		s stable for 24 months from da	ate of manufacture. ad, performance must be validated by the user.
Procedure:			
1.	Tissue Section Pretreatment: Staining of formalin fixed, paraffin embedded tissue sections is enhanced by pretreatment with Citrate Plus (ScyTek catalog# CPL500) or 10mM citrate buffer, pH 6.0 (ScyTek Catalog# CBB500, see IFU for instructions). Enzymatic predigestion with Pepsin (ScyTek catalog# PSS060, see IFU for instructions) is required prior to staining.		
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 "Retrieval HRP Anti-Polyvalent Lab Pack" (ScyTek catalog# RPL125, see IFU for instructions) combined with the "DAB Chromogen/Substrate Bulk Pack (High Contrast)" (ScyTek catalog# ACV500, see IFU for instructions). 		
Storage: 2°C	8°C	ScyTek Laboratories, In 205 South 600 West Logan, UT 84321 U.S.A.	nc. IVD Ec REP EmergoEurope (31)(0) 70 345-8570 Molsnstraat 15 2513 BH Hague

2513 BH Hague The Netherlands

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LaboratoriesInstructions For Use
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Precautions: Contains Sodium Azide as a preservative (0.09% w/v).

Do not pipette by mouth.

Avoid contact of reagents and specimens with skin and mucous membranes. Avoid microbial contamination of reagents or increased nonspecific staining may occur. This product contains no hazardous material at a <u>reportable concentration</u> according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC.

References:

- 1. Autio-Harmainen H, et al. Synthesis of laminin and type IV collagen by trophoblastic cells and fibroblastic stromal cells in the early human placenta. Lab Invest 1991; 64:483.
- 2. Engval E, et al. Distribution and isolation of four laminin variants; tissue restricted distribution of heterodimers assembled from five different subunits. Cell Regulation 1990; 1:731.
- 3. Klenman HK, et al. Laminin receptors for neurite formation. Proc Natl Acad Sci USA 1988; 85:1282.
- Warranty: No products or "Instructions For Use (IFU)" are to be construed as a recommendation for use in violation of any patents. We make no representations, warranties or assurances as to the accuracy or completeness of information provided on our IFU or website. Our warranty is limited to the actual price paid for the product. ScyTek Laboratories, Inc. is not liable for any property damage, personal injury, time or effort or economic loss caused by our products. 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.

