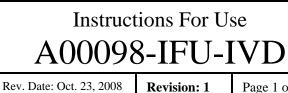
ScyTek Laboratories



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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

## Cytokeratin (Pan)

| Availability/Co   | ontents:  | <u>Item #</u><br>A20098<br>A00098<br>A00098.0025  | <u>Volume</u><br>2 ml<br>6 ml<br>25 ml |  |
|---|---|---|--|--|
| Description:  |   |   |  |  |
| Species:<br>Immunogen:<br>Clone:<br>Isotype:<br>Format:   |   | Mouse<br>Multiple<br>Multiple<br>IgG<br>This antibody has been pretitered and quality controlled to work on formalin-fixed paraffin-<br>embedded as well as acetone fixed cryostat tissue sections. No further titration is required. |  |  |
| Specificity:  |   | This antibody cocktail reacts with keratins 4, 5, 6, 7, 8, 10, 13, 14, 15, 16, 18, and 19.<br>Cytokeratin (Pan) differentiates epithelial tumors from non-epithelial tumors.  |  |  |
| Species Reactivity:<br>Positive Control:<br>Cellular Localization:<br>Titer/Working Dilution:<br>Microbiological State: |   | Human, Pig. Others not test<br>Skin<br>Cytoplasmic<br>No further dilution is required<br>This product is not sterile.   | ed.                                    |  |
| This proc<br>fixed, pa  |   | tro Diagnostic Use.<br>duct is intended for qualitative immunohistochemistry with normal and neoplastic formalin-<br>raffin-embedded tissue sections, to be viewed by light microscopy.<br>Ise past expiration date.                  |  |  |
|   |   | ntigrade.<br>is stable for 24 months from date of manufacture.<br>it is not stored as recommended, performance must be validated by the user.   |  |  |
| Procedure:  |   |   |  |  |
| 1.  | Tissue Section Pretreatment: Staining of formalin fixed, paraffin embedded tissue sections requires pretreatment Trypsin, Stabilized Solution (ScyTek catalog# TSS155) for 10 minutes at 37° C prior to staining.   |   |  |  |
| 2.  | Primary Antibody Incubation Time: We suggest an incubation period of 30 minutes at room temperature.<br>However, depending upon the fixation conditions and the staining system employed, optimal incubation should<br>be determined by the user.   |   |  |  |
| 3.  | <ol> <li>Visualization: For maximum staining intensity we recommend the "UltraTek HRP Anti-Polyvalent Lab Pack"<br/>(ScyTek catalog# UHP125, see IFU for instructions) combined with the "DAB Chromogen/Substrate Bulk Pack<br/>(High Contrast)" (ScyTek catalog# ACV500, see IFU for instructions).</li> </ol> |   |  |  |
| ŀ   | 8°C   | ScyTek Laboratories, I<br>205 South 600 West  | nc.                                    |  |

Storage: 2°C

205 South 600 West Logan, UT 84321 U.S.A.

EmergoEurope (31)(0) 70 345-8570 Molsnstraat 15 2513 BH Hague The Netherlands

## ScyTek<br/>LaboratoriesInstructions For Use<br/>A00098-IFU-IVDRev. Date: Oct. 23, 2008Revision: 1Page 2 of 2

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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. Woodcock-Mitchell J., et al. Journal of Cell Biology, Nov., 95 (2 Pt 1): pages 580-588, 1982.
- 2. Tseng S.C., et al. Cell, Sept., 30(2): pages 361-372, 1982.
- 3. Eichner R., et al. Journal of Cell Biology, April, 98(4): pages 1388-1396, 1984.
- 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.



