

Instructions For Use			
RA0078-C.5-IFU-RUO			
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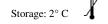
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p27^{Kip1} (Mitotic Inhibitor/Suppressor Protein); Clone KIP1/769 (Concentrate)

Availability/Contents: Description:	Item # Volume RA0078-C.5 0.5 ml	
Species: Immunogen: Clone: Isotype: Entrez Gene ID:	Mouse Recombinant human CDKN1B protein KIP1/769 IgG1, kappa 1027 (Human)	
Hu Chromosome Loc.: Synonyms: Mol. Weight of Antigen:	12p13.1 CDKN1B, CDKN4, Cyclin Dependent Kinase Inhibitor 1B, Cyclin-dependent kinase inhibitor p27 Kip1, KIP1, MEN1B, MEN4 25-26kDa	
Format: Specificity: Background:	200μg/ml of Ab purified from Bioreactor Concentrate by Protein A/G. Prepared in 10mM PBS with 0.05% BSA & 0.05% azide. This antibody recognizes a 27kDa protein, identified as p27 ^{Kip1} , a cell cycle regulatory mitotic inhibitor. It is highly specific and shows no cross-reaction with other related mitotic inhibitors. p27 ^{Kip1} functions as a negative regulator of G1 progression and has been proposed to function as a possible mediator of TGF-β induced G1 arrest. p27 ^{Kip1} is a candidate tumor suppressor	
Species Reactivity: Positive Control: Cellular Localization: Titer/ Working Dilution:	 gene. This MAb is excellent for staining of formalin-fixed tissues. Human, Mouse, Rat and Monkey. Others not known. ZR75, T47D, SK-BR-3, MDA-MB-231, MCF7 cells. Tonsil, breast or colon carcinoma. Nuclear Immunohistochemistry (Frozen and Formalin-fixed): 0.5-1 μg/ml Flow Cytometry: 0.5-1 μg/million cells Immunofluorescence: 0.5-1 μg/ml 	
Microbiological State:	Western Blotting:0.5-1 µg/mlImmunoprecipitation:0.5-1 µg/500µg protein lysateThis product is not sterile.	





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Ordering Information and Current Pricing at www.scytek.com

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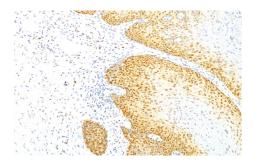
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Uses/Limitations:

Not to be taken internally. For Research Use Only. 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 if reagent becomes cloudy. Do not use past expiration date. Non-Sterile.



Formalin-paraffin human cervical cancer stained with p27^{Kip1}; Clone KIP1/769.

Procedure:

- 1. **Tissue Section Pretreatment (Required):** Staining of formalin fixed, paraffin embedded tissue sections is significantly enhanced by pretreatment with Citrate Plus (ScyTek catalog# CPL500).
- 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 "UltraTek HRP Anti-Polyvalent Lab Pack" (ScyTek catalog# UHP125, see IFU for instructions) combined with the "DAB Chromogen/Substrate Bulk Pack (High Contrast)" (ScyTek catalog# ACV500, see IFU for instructions).

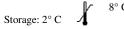
 Precautions:
 Contains Sodium Azide as a preservative (0.09% w/v).

 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 reportable concentration according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC.

References:

- 1. Fredersdorf S, Burns J, Milne AM, Packham G, Fallis L, Gillett CE, et al. High level expression of p27Kip1and cyclin D1 in some human breast cancer cells: Inverse correlation between the expression of p27Kip1 and degree of malignancy in human breast and colorectal cancers. Proc Natl Acad Sci 1997;94:6380-5.
- 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.





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