

# Produktinformation



Forschungsprodukte & Biochemikalien
Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart siehe unsere Liefer- und Versandbedingungen

## Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

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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

# CD57 (HNK-1); Clone NK-1 (Ready-To-Use)

Availability/Contents:	<u>Item #</u> A00117-0002 A00117-0007 A00117-0025	<u>Volume</u> 2 ml 7 ml 25 ml
Description:		
Species: Immunogen: Clone: Isotype: Format: Specificity: Background:	NK-1 Mouse IgM, Kappa This antibody has been p embedded as well as acc The NK1 antibody clone CD57 is a terminally su HNK (human natural kille CD stands for cluster of term CD57 has been u lymphocytic subpopulation cells and are commonly tonsils. CD57 antibody p replicative senescence (of susceptibility to activati positivity patterns have b and malignancies. Increa allogenic transplantation, It is increasingly being re most often referred to as peripheral nerve tissue w and neurogenic patholog to identify neuroendocrim that HNK1 autoantibodies	mononuclear cells were used as immunogen. pretitered and quality controlled to work on formalin-fixed paraffin- etone fixed cryostat tissue sections. No further titration is required. recognizes the glycoepitope referred to as both CD57 and HNK1. Ifated glycan carbohydrate epitope (glycoepitope) first discovered on er) cells in 1981. CD57 is also referred to as CD57 antigen and HNK1. f differentiation and HNK1 for human natural killer1. Historically, the sed in immunology and antibody to CD57 is important for defining ons. CD57 antibody positive lymphocytes are typically either T or NK r found within the germinal centers of the spleen, lymph nodes and positivity in T lymphocytes has long been used as a marker of in vitro clonal exhaustion). CD57 antibody positive T lymphocytes have a high on-induced death. CD57 upregulation or unusual CD57 antibody poen identified in diseases including autoimmunity, chronic infections, pased CD57 antibody positivity has also been associated with aging, and even physical and psychological stress. cognized that CD57 has important roles in the nervous system where it HNK1. HNK1 (CD57) is predominantly expressed in brain and there it is involved in development, homeostasis, normal development y. For example, HNK1 antibody positivity has been used as a marker e cells and their tumors as expression is high in both. It is also notable s have been detected in peripheral demyelinating neuropathy ince of the immune system in neurological function.
Species Reactivity: Positive Control: Cellular Localization: Titer/Working Dilution: Microbiological State:	Human. Tonsil, spleen, lymph noo Membrane. No further dilution is requ This product is not sterile	iired.

Storage: 2° C



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## CE

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

Doc: IFU-Template2-8rev2



## Instructions For Use A00117-IFU-RUO

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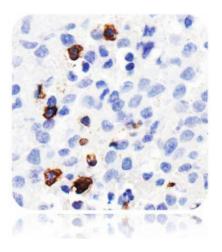
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**Uses/Limitations:** 

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. Use caution when handling reagents. Non-Sterile.

### Ordering Information and Current Pricing at www.scytek.com



### **Procedure:**

- 1. **Tissue Section Pretreatment:** Staining of formalin fixed, paraffin embedded tissue sections is enhanced by pretreatment with Citrate Plus (ScyTek catalog# CPL500) or Citrate Buffer (10x), pH 6.0 (ScyTek Catalog# CBB500, see IFU for instructions).
- 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).

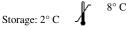
Precautions: Contains Sodium Azide as a preservative (0.09% w/v). Do not pipette by mouth. Avoid contact of reagents and specimens with skin and mucous membranes. Avoid microbial contamination of reagents or increased nonspecific staining may occur. This product contains no hazardous material at a <u>reportable concentration</u> according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC.

### **References:**

- 1. Burger D, AJ Steck, CC Bernard, N Kerlero de Rosbo. Journal Neurochem 61:1822-1827 (1993).
- 2. Guarino M. Pathol Res Pract 189:913-920 (1993).
- 3. Cavazzana AO, V Ninfo, J Roberts, TJ Triche. Modern Pathol 5:71-78 (1992).
- 4. Focosi D, M Bestagno, O Burrone, M Petrini. J Leukoc Biol. 87:107-116 (2010).
- 5. Kizuka Y, S Oka. Cell Mol Life Sci. DOI 10.1007/s00018-012-1036-z (2012).

### 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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