

Instructions For Use

A00022-IFU-IVD

Rev. Date: Jan. 8, 2024 Revision: 3

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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

PCNA; Clone PC10

Catalog Number	<u>Format</u>	Volume
A00022-0002	(Ready-To-Use)	2 ml
A00022-0007	(Ready-To-Use)	7 ml
A00022-0025	(Ready-To-Use)	25 ml
A00022-C.1	(Concentrate)	0.1 ml
A00022-C	(Concentrate)	1 ml

Intended Use

For In-Vitro Diagnostic Use. 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.

Description

Titer/Working Dilution: Ready-to-Use: No further dilution required.

Concentrate: Suggested dilution is 1:100-200

Species: Mouse

Immunogen: Rat PCNA/Protein A fusion protein.

Clone: PC10 lgG2a, kappa

Entrez Gene ID: 5111 (Human); 18538 (Mouse); 25737 (Rat)

Hu Chromosome Loc.: 20p12.3

Synonyms: Cyclin; DNA polymerase delta auxiliary protein; Mutagen-

sensitive 209 protein; PCNAR; Polymerase delta accessory

protein

Mol. Wt. of Antigen: 36 kDa

Format: Ready-To-Use 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.

Concentrate antibody is provided at 200µg/ml of Ab purified from Bioreactor Concentrate by Protein A/G. Prepared in 10mM PBS

with 0.05% BSA & 0.05% Sodium Azide.

Specificity: Recognizes a non-histone protein of 36kDa, which is identified as

proliferating cell nuclear antigen (PCNA). It is also known as

cyclin or polymerase delta auxiliary protein.

Background: Elevated expression of PCNA/cyclin has been shown in the

nucleus during late G1 phase immediately before the onset of DNA synthesis, becoming maximal during S-phase, and declining during G2 and M phases. This antibody is excellent for multiple

applications.

Species Reactivity: Human, Monkey, Pig, Mouse, Rat, Chicken, Zebrafish,

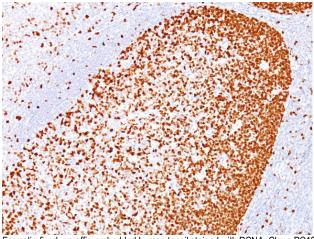
Drosophila melanogaster and Yeast (S. pombe & S. cerevisiae).

Others-not known.

Positive Control: Tonsil or reactive lymph node.

Cellular Localization: Predominantly nuclear, some cytoplasmic

Microbiological State: Nonsterile



Formalin-fixed, paraffin-embedded human tonsil stained with PCNA; Clone PC10.

Materials and Reagents Required but not Provided

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

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

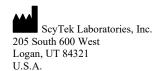
Procedure

- 1. Tissue Section Pretreatment (Highly Recommended): Staining of formalin fixed paraffin embedded tissue sections is significantly enhanced by pretreatment with Tis-EDTA HIER Solution (10x) pH 9.0 (ScyTek catalog# TES500) or Citrate Plus (10x) HIER Solution (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 "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).

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.







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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 <u>reportable concentration</u> according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC.
- 2. Do not pipette by mouth.
- 3. Avoid contact of reagents and specimens with skin and mucous membranes.
- 4. Avoid microbial contamination of reagents or increased nonspecific staining may occur.
- 5. The user must validate any procedures and recommendations that differ from this data sheet.
- 6. The SDS may be found at scytek.com

References

- Waseem NH & Lane DP. 1990. J Cell Sci. 96:121-9.
- 2. Hall PA et al. 1990. J. Pathol. 162(4):285-94.
- 3. Landberg G & Roos G. 1991. Cancer Res. 51 (17):4570-4.
- Woods AL et al. 1991. Histopathol. 19(1):21-7
- 5. Yu, CC et al. 1991. Histopathol. 19(1):29-33.

Warranty

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