

CD4 (Clone: EP204)

**Rabbit Monoclonal Antibody** 

PRODUCT INFORMATION:

PERFORMANCE CHARACTERISTICS:

REF PR013 6ml Ready to use PR013

Localization: Membrane 3ml Ready to use **CR013** 1ml Concentrated **CR013** 0.5ml Concentrated

**CR013** 0.1ml Concentrated 6ml Ready to use **HAR013** HAR013 3ml Ready to use

Retrieval Buffer: EDTA, pH 8.0 Incubation: 60 minutes

Positive control: Tonsil, T-Cell Lymphoma

### INTENDED USE

#### For in vitro diagnostic use only

This antibody is intended for use in qualitatively identifying CD4 antigen by light microscopy in formalin fixed, paraffin embedded (FFPE) tissue sections using immunohistochemical (IHC) detection methodology. Interpretation of any positive or negative staining must be complemented with the evaluation of proper known controls (Positive and Negative) and must be made within the context of the patient's clinical history and other diagnostic tests. A qualified and trained pathologist must perform the evaluation of the test. This antibody is intended to be used after the primary diagnosis of tumor has been made by conventional histopathology using non-immunologic histochemical stains.

#### **SUMMARY AND EXPLANATION**

CD4 is a glycoprotein found on the surface of immune cells such as T helper cells, monocytes, macrophages and dendritic cells. It is a co-receptor that assists the Tcell receptor (TCR) with an antigen- presenting cell and also interacts directly with MHC class II molecules on the surface of the antigen-presenting cells using its extracellular domain. In lymphatic tissues, the CD4+ T-cells are seen in large numbers in the parafollicular zone, while scattered cells are found in the germinal centers and mantle zone. CD4 is also demonstrated in hepatic sinusoidal cells, monocytes and monocytes-derived cells but not expressed on B-cells and immature thymocytes. Precursor T- lymphoblastic lymphomas are therefore variable in their expression of CD4. Most mature T-cell lymphomas are CD4 positive with the exception of aggressive NK-cell leukemia and extra nodal NK/T-cell lymphoma.

CD4 plays an important role in the classification of lymphocytes in inflammatory lesions and malignant lymphomas.

## PRINCIPLE OF THE PROCEDURE

The identification of the antigen on the FFPE tissues is carried out using the above stated antibody. The antigen and antibody complex are visualized using an enzyme coupled (HRP/AP) secondary antibody with specific binding to the primary antibody, this complex is visualized by the enzymatic activation of the chromogen resulting in a visible reaction production of the antigenic site. Each and every step involves precise time and optimal temperature and the results are interpreted using a light microscope by a qualified and trained pathologist.

## REAGENT PROVIDED

Concentrated format: CD4 antibody is affinity purified and diluted in antibody diluent with 1% bovine serum albumin (BSA) and 0.05% sodium azide (NaN3). Recommended dilutions: 1:50 - 1:100

The antibody dilution and protocol may vary depending on the specimen preparation and specific application. Optimal conditions should be determined by individual laboratories.

Pre-diluted format: PathnSitu's ready to use antibodies are pre-tittered to optimal staining conditions. Further dilution will affect the efficacy of the antibody and may yield to sub-optimal staining.

Immunogen: A synthetic peptide corresponding to residues of human CD4 protein

Host, Isotype: Rabbit, IgG

DS-PR013-B.

#### STORAGE AND HANDLING

Storage Recommendations: Store at 2-8°C. When stored at the appropriate conditions, the antibody is stable until expiry. Do not use the antibody after the expiration date provided on the vial in any condition.

To ensure proper reagent delivery and stability, replace the dispenser cap after every use and immediately place the vial into the refrigerated conditions in an

upright position. The contents of the vial should be used within 9 months from the opening of the vial.

#### SPECIMEN PREPARATION

#### Staining Recommendations:

Routinely processed, FFPE tissues are suitable for use with this primary antibody. when using PathnSitu's Poly Excel HRP/DAB detection system. The recommended tissue fixative is 10% neutral buffered formalin. Variable results may occur as a result of prolonged fixation or special processes such as decalcification. The thickness of the sections should be 2-5 µm. Slides should be stained once the sections are made as the antigenicity of the cut sections may diminish over a period of time. It is recommended to stain known positive and negative controls simultaneously with unknown specimens.

### **PRECAUTIONS**

- This product should be used by qualified and trained professional users only
- The product contains < 0.1% of sodium azide as a preservative and is not classified as hazardous, refer to MSDS for further details
- 3. As with any product derived from biological sources, proper handling procedures should be used
- 4. Do not use reagents after the expiration date
- 5. Use protective clothing and gloves, while handling reagents
- All hazardous materials should be disposed of according to local state and federal regulations
- Avoid microbial contamination of reagents as it may lead to incorrect results

## STAINING PROCEDURE

Antigen Retrieval Solution: Use EDTA Buffer (Cat#PS008) as an antigen retrieval solution.

Heat Retrieval Method: Retrieve sections under steam pressure for 20 minutes using PathnSitu's MERS (Multi Epitope Retrieval System) for optimal retrieval of the epitopes, allow solution to cool at room temperature, transfer the tissue sections/slides to the distilled water prior to the primary antibody application.

Primary Antibody: Cover the tissue sections with primary antibody and incubate for 60 min at room temperature when used PathnSitu's PolyExcel Detection System.

Detection System: Refer to PathnSitu's PolyExcel HRP/ DAB detection system protocol for optimal staining results.

## **QUALITY CONTROL**

The recommended positive tissue control for CD4 are Tonsil and T Cell Lymphoma. A positive and negative tissue controls must be run with every staining procedure performed for monitoring the correct performance of processed tissue and test reagents. A negative tissue control provides an indication of non-specific background staining. If the results are not expected in positive and negative controls the test must be considered invalid and the entire procedure must be cross verified. The individual laboratory must establish their own quality control to validate the process and antibody when opening a vial.

## INTERPRETATION OF RESULTS

CD4 stains the Membrane. A qualified experienced/trained pathologist must interpret the results in the patient's sample along with the positive and negative controls

# PERFORMANCE CHARACTERISTICS

PathnSitu products will undergo thorough a quality control check before it is released to the market. The antibody showed consistent specific and sensitive staining on the multiple positive tissue controls tested, by inter run, intra run and lot based studies. The antibody is stable for the expiry mentioned on the labels which is determined by real time or accelerated methods.

### TROUBLESHOOTING

- Follow the antibody specific protocol recommendations according to the data sheet provided
- Tissue staining is dependent on the handling and processing of the tissue prior to staining. Improper fixation, tissue processing, antibody freezing and thawing, washing, drying, heating, sectioning or contamination with other

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- tissues or fluids may produce artifacts, antibody trapping or inaccurate results
- 3. Do not allow the section to dry out during the entire IHC process
- Excessive or incomplete counter staining may compromise the interpretation of the results
- If unusual results occur, contact PathnSitu's Technical Support at +91-40-2701 5544 or E-mail:techsupport@pathnsitu.com

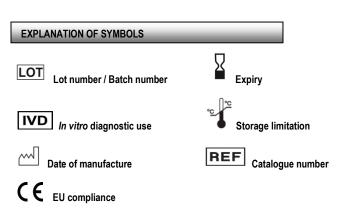
#### LIMITATIONS AND WARRANTY

Authorized and skilled/trained personnel only may use the product. The clinical interpretation of any test results should be evaluated within the context of the patient's medical history and other diagnostic test results. A qualified trained pathologist must perform the evaluation of the test results. There are no warranties, expressed or implied, which extend beyond the description. PathnSitu is not liable for property damage, personal injury, time or effort on economic loss caused by this product.

## **BIBLIOGRAPHY**

- 1. Brady RL, et al.: Science 1993, 260:979-983
- 2. Sattentau QJ, et al.: Cell 1988, 52:631-633
- 3. Williamson SL, et al.: Am J Pathol 1998, 152:1421-1426
- Dabbs DJ. Diagnostic Immunohistochemistry 2010; Churchill Livingstone

CD4, EP204 antibody has been created by Epitomics Inc., using Epitomics proprietary rabbit monoclonal antibody technology covered under Patent No.'s 5,675,063 and 7,402,409.



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