

# Neurofilament Primary Antibody

**Clone: NE14 (Mouse Polyclonal)**

**Localization** – Cytoplasm

**Host Species** – Rabbit

**Ig Class** – IgG1 / Kappa

## Intended Use

This antibody is designed for the specific localization of Neurofilament in formalin-fixed, paraffin-embedded (FFPE) tissue sections.

## Storage & Handling

Store RTU Vial at 2-8°C. Fresh dilutions for concentrated antibodies, if required, should be prepared prior to use and are stable for up to one day at room temperature (20-26°C).

## Working Principle

IHC is a two-step process wherein the primary antibody binds to the antigen of interest and that binding is detected by a chromogen. The primary antibody may be used in IHC using manual techniques or any Automated Staining System. Positive and negative controls should always be run simultaneously with all patient specimens.

## Product Description

This monoclonal antibody recognizes a 200 kDa protein corresponding to the heavy subunit of neurofilaments (NF-H). Its reactivity is specific to the phosphorylated KSP/KEP repeat domain located at the C-terminal region of NF-H. Following dephosphorylation of neurofilaments with alkaline phosphatase, antibody binding is abrogated, confirming phosphorylation-dependent recognition. Neurofilaments constitute the principal structural scaffolding of axons and dendrites and are distributed throughout neurons, peripheral nerves, and sympathetic ganglion cells. They are composed of three major subunits: NF-L (68 kDa), NF-M (160 kDa), and NF-H (200 kDa). Immunoreactivity with anti-neurofilament is observed across a spectrum of neural, neuroendocrine, and endocrine neoplasms. Positive staining is typically seen in neuromas, ganglioneuromas, gangliogliomas, ganglioneuroblastomas, and neuroblastomas. Additionally, neurofilament expression is detected in paragangliomas, as well as in adrenal and extra-adrenal pheochromocytomas. Carcinoid tumors, cutaneous neuroendocrine carcinomas, and pulmonary small-cell (oat cell) carcinomas also demonstrate immunopositivity for neurofilament.

## Material Supplied

Neurofilament antibody is affinity purified and diluted in PBS, pH 7.4, containing 1% BSA and 0.09% sodium azide.

## Material required But Not Supplied

- Xylene
- DI Water
- Control Tissues
- Isopropyl alcohol
- Antigen retrieval buffers
- Hematoxylin
- Positive charged slides
- Blocking Reagents
- Mounting media
- Wash Buffer
- Detection System
- Cover glass

## Working Reagent Procedure

- Ready-to-Use antibodies have been optimized for use with the recommended protocols and should not require further dilution.
- Concentrated antibodies must be diluted in accordance with the recommended protocol.

## Recommended Protocol

Refer the following table for the details on specific recommended protocol for this antibody.

|                        |   |                                 |  |
|------------------------|---|---------------------------------|--|
| <b>Control Tissue</b>  | Neuroblastoma                               | <b>Antibody Incubation Time</b> | 30-60 Minutes at RT                                |
| <b>Dilution factor</b> | <b>1:20-50</b><br>(Antibody Diluent: DH144) | <b>Retrieval Pre-treatment</b>  | <b>Tris-EDTA based HIER</b><br>(AR9 Buffer: DH020) |

## Precautions

*This product should be used by qualified and trained professional users only.*

Avoid microbial contamination of reagents to minimize non-specific staining. Never pipette reagents by mouth. Avoid contact of reagents and specimens with skin. If reagents or specimens come into contact with sensitive area, wash with sufficient amounts of water. Dispose of the unused reagents. This kit contain sodium azide at concentrations of less than 0.1%. Sodium azide is not classified as a hazardous chemical at these concentrations, but proper handling protocols should be observed. For more information on product hazards, precautions and waste disposal, *Material Safety Data Sheets* are available upon request.










## Limitations

Improper tissue handling and processing prior to immunostaining can lead to inconsistent results. Variations in embedding and fixation or the nature of the tissue may lead to variations in results. Endogenous peroxidase activity or pseudo peroxidase activity in erythrocytes and tissue biotin may result in non-specific staining based on the detection system employed. Tissues containing Hepatitis B Surface Antigen (HBsAg) may give false positive with horseradish peroxidase systems. Improper counterstaining and mounting may compromise the interpretation of results. Interpretation of the staining result is solely the responsibility of the user. Experimental results should be confirmed by a medically-established diagnostic product or procedure. Evaluation must be performed by a qualified pathologist.

## Troubleshooting

For Technical Support contact us at +91 - 7506501122 or [info@dygnova.com](mailto:info@dygnova.com) or your local distributor to report unusual staining.

Doc No: DH/DS/PR1535Rev.00

|   |                      |   |                                   |   |                                    |
|---|----------------------|---|-----------------------------------|---|------------------------------------|
|  | Manufacturer Details |  | Use by Date                       |  | Lot/Batch Number                   |
|  | Manufacturing Date   |  | Consult Instructions for Use      |  | Catalogue Number                   |
|  | Temperature Limits   |  | Sufficient for 'n' assays / tests |  | In-vitro Diagnostic Medical Device |

