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# **Product Information**

# **Anti-Neurofilament 200-FITC Antibody**

Mouse monoclonal, Clone NE14, purified from hybridoma cell culture

Product Number SAB4200811

## **Product Description**

Monoclonal Anti-Neurofilament 200-FITC (mouse IgG1 isotype) is derived from the NE14 hybridoma produced by the fusion of mouse myeloma cells and splenocytes from an immunized mouse. Neurofilaments purified from pig spinal cord were used as the immunogen. The isotype is determined by ELISA using Mouse Monoclonal Antibody Isotyping Reagents (Product Number ISO2). The antibody is purified from culture supernatant of hybridoma cells. The purified antibody is conjugated to fluorescein isothiocyanate (FITC).

Monoclonal Anti-Neurofilament 200, also known as Neurofilament-H or Heavy subunit, specifically recognizes the phosphorylated H tail of Neurofilament 200 and shows no reactivity toward enzymatically dephosphorylated neurofilaments. The antibody shows reactivity with neurofilaments in the central and peripheral nervous systems from human, pig, mouse, rat, chicken, guinea pig, feline, and bovine origin. The antibody may be used in various immunochemical techniques including immunohistochemistry and immunoblotting (~200 kDa).

Neurofilaments are a type of intermediate filaments (IFs) that serve as major elements of the cytoskeleton supporting the axon cytoplasm of neuronal cells. IFs are components of most eukaryotic cells and significantly differ from other cytoskeletal elements of the cell, namely microtubules and microfilaments.

Neurofilaments undergo post-translational modifications including different levels of phosphorylation, which has been suggested to modulate their function by influencing the interaction between neurofilament and cytoplasmic organelles. Neurofilaments are built from three intertwined protofibrils of apparent molecular masses [68 (L), 160 (M), and 200 (H) kDa] which are themselves composed of two tetrameric protofilament complexes of monomeric proteins. Neurofilament 200 also known as Neurofilament heavy polypeptide (H-subunit), NF-H, NEFH or 200 kDa neurofilament protein, has an important function in mature axons that is not subserved by the two smaller neurofilament proteins.

Defects in Neurofilament 200 are a cause of susceptibility to amyotrophic lateral sclerosis (ALS) and these accumulations are a hallmark of pathological lesions. Begin Neurofilaments can accumulate in large numbers within cell bodies and proximal axons of affected neurons in several pathological diseases, such as Charcot-Marie-Tooth (CMT), neurofilament inclusion disease (NFID), giant axonal neuropathy (GAN), diabetic neuropathy, spinal muscular atrophy (SMA) and spastic paraplegia. In addition, neurofilament accumulations was detected in Alzheimer's (AD) and Parkinson's disease (PD) patients.

# Reagent

Supplied as a solution in 0.01 M phosphate buffered saline, pH 7.4, containing 15 mM sodium azide as a preservative.

Antibody Concentration: ~1.0 mg/mL

## **Precautions and Disclaimer**

For R&D use only. Not for drug, household, or other uses. Please consult the Safety Data Sheet for information regarding hazards and safe handling practices.

# Storage/Stability

For continuous use, store at 2—8 °C for up to one month. For extended storage, freeze in working aliquots. Repeated freezing and thawing is not recommended. If slight turbidity occurs upon prolonged storage, clarify the solution by centrifugation before use. Working dilution samples should be discarded if not used within 12 hours. Protect from prolonged exposure to light.

#### **Product Profile**

Direct Immunohistochemistry: a working dilution of 1:200—1:400 (4-8 μg/mL) is recommended using enzyme treated formalin-fixed, paraffin-embedded rat cerebellum sections.

<u>Note</u>: In order to obtain best results in different techniques and preparations, it is recommended to determine optimal working concentration by titration.

#### References

- 1. Debus, E. et al., *Differentiation*, **25**, 193-203 (1983).
- 2. Shaw, G. et al., Eur. J. Cel Biol., 42, 1-9 (1986).
- 3. Shaw, G. et al., Eur. J. Cell Biol., 34, 130-6 (1984).
- 4. Dahl, D., J. Neurosci. Res., 20, 431-41 (1988).
- 5. Saito, Y. et al.. eLife, 5, e14371 (2016).
- 6. Jacques, T.S. et al., EMBO, 29, 222-35 (2010).
- Bocquet, A. et al., J. Neurosci., 29, 11043-54 (2009).
- 8. Lu, C.H. et al., PLoS One, 7, e40998 (2012).
- 9. Yuan, A. et al., J. Cell Sci., 125, 3257-63 (2012).
- 10. Abe, A. et al., J. Hum. Genet., 54, 94-7 (2009).
- 11. Liu, Y.L. et al., Spinal Cord, 47, 166-70 (2009).
- 12. Szaro, B.G. et al., *Trends Neurosci.*, **33**, 27–37 (2010).

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