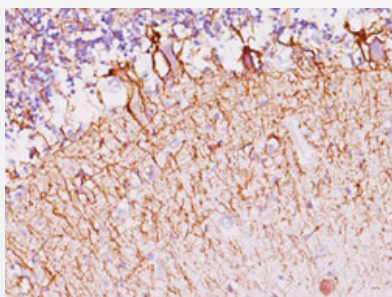


NEFH/NEFL monoclonal antibody, clone SPM145

Catalog # MAB13212 Size 100 ug

Applications



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining (Formalin-fixed paraffin-embedded sections) of human cerebellum with NEFH/NEFL monoclonal antibody, clone SPM145 (Cat # MAB13212).

Specification

Product Description	Mouse monoclonal antibody raised against native human NEFH/NEFL.
Immunogen	Semi-purified NEFH/NEFL from human spinal cord.
Host	Mouse
Theoretical MW (kDa)	200, 68
Reactivity	Human
Form	Liquid
Purification	Protein G purification
Isotype	IgG1
Recommend Usage	Flow Cytometry (0.5-1 ug/10 ⁶ cells in 0.1 mL) Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) (0.25-0.5 ug/mL) The optimal working dilution should be determined by the end user.
Storage Buffer	In 10 mM PBS.

Storage Instruction

Store at -20 to -80°C.
Aliquot to avoid repeated freezing and thawing.

Applications

- Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Immunohistochemical staining (Formalin-fixed paraffin-embedded sections) of human cerebellum with NEFH/NEFL monoclonal antibody, clone SPM145 (Cat # MAB13212).

- Flow Cytometry

Gene Info — NEFH

Entrez GeneID [4744](#)

Protein Accession# [P12036;P07196](#)

Gene Name NEFH

Gene Alias NFH

Gene Description neurofilament, heavy polypeptide

Omim ID [105400 162230](#)

Gene Ontology [Hyperlink](#)

Gene Summary Neurofilaments are type IV intermediate filament heteropolymers composed of light, medium, and heavy chains. Neurofilaments comprise the axoskeleton and functionally maintain neuronal caliber . They may also play a role in intracellular transport to axons and dendrites. This gene encodes the heavy neurofilament protein. This protein is commonly used as a biomarker of neuronal damage and susceptibility to amyotrophic lateral sclerosis (ALS) has been associated with mutations in this gene. [provided by RefSeq]

Other Designations neurofilament triplet H protein|neurofilament, heavy polypeptide 200kDa

Gene Info — NEFL

Entrez GeneID [4747](#)

Protein Accession# [P12036;P07196](#)

Gene Name NEFL

Gene Alias	CMT1F, CMT2E, FLJ53642, NF-L, NF68, NFL
Gene Description	neurofilament, light polypeptide
Omim ID	162280 607684 607734
Gene Ontology	Hyperlink
Gene Summary	Neurofilaments are type IV intermediate filament heteropolymers composed of light, medium, and heavy chains. Neurofilaments comprise the axoskeleton and they functionally maintain the neuronal caliber. They may also play a role in intracellular transport to axons and dendrites. This gene encodes the light chain neurofilament protein. Mutations in this gene cause Charcot-Marie-Tooth disease types 1F (CMT1F) and 2E (CMT2E), disorders of the peripheral nervous system that are characterized by distinct neuropathies. A pseudogene has been identified on chromosome Y. [provided by RefSeq]
Other Designations	light molecular weight neurofilament protein neurofilament protein, light chain neurofilament subunit NF-L neurofilament triplet L protein neurofilament, light polypeptide 68kDa neurofilament-light

Pathway

- [Amyotrophic lateral sclerosis \(ALS\)](#)
- [Amyotrophic lateral sclerosis \(ALS\)](#)

Disease

- [Amyotrophic lateral sclerosis](#)
- [Dominance](#)
- [Genetic Predisposition to Disease](#)
- [Genetic Predisposition to Disease](#)
- [Kidney Failure](#)
- [Motor Neuron Disease](#)
- [Multiple Sclerosis](#)
- [Multiple Sclerosis](#)
- [Parkinson disease](#)
- [Schizophrenia](#)