

# **Product datasheet for CF814222**

#### OriGene Technologies, Inc.

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## Neurofilament (NEFL) Mouse Monoclonal Antibody [Clone ID: OTI3F8]

### **Product data:**

**Product Type:** Primary Antibodies

Clone Name: OTI3F8
Applications: ELISA

Recommended Dilution: ELISA 1:5000-10000

Reactivity: Human
Host: Mouse
Isotype: IgG1

Clonality: Monoclonal

**Immunogen:** Human recombinant protein fragment corresponding to amino acids 401-543aa of human

NEFL (NP\_006149) produced in E.coli.

Formulation: Lyophilized powder (original buffer 1X PBS, pH 7.3, 8% trehalose)

**Reconstitution Method:** For reconstitution, we recommend adding 100uL distilled water to a final antibody

concentration of about 1 mg/mL. To use this carrier-free antibody for conjugation experiment, we strongly recommend performing another round of desalting process. (OriGene recommends Zeba Spin Desalting Columns, 7KMWCO from Thermo Scientific)

**Purification:** Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography

(protein A/G)

Conjugation: Unconjugated

**Predicted Protein Size:** 61.5 kDa

**Gene Name:** neurofilament light chain

Database Link: NP 006149

Entrez Gene 4747 Human

P07196





### Neurofilament (NEFL) Mouse Monoclonal Antibody [Clone ID: OTI3F8] - CF814222

**Background:** 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, Oct 2008]

Synonyms: CMT1F; CMT2E; CMTDIG; NF-L; NF68; NFL; PPP1R110

Protein Families: Druggable Genome, ES Cell Differentiation/IPS

**Protein Pathways:** Amyotrophic lateral sclerosis (ALS)