

Product datasheet for RC205920L2V

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Neurofilament (NEFL) (NM_006158) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Neurofilament (NEFL) (NM_006158) Human Tagged ORF Clone Lentiviral Particle

Symbol: Neurofilament

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

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_006158 **ORF Size:** 1629 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC205920).

Sequence:

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of

reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

RefSeg: NM 006158.2

 RefSeq Size:
 3854 bp

 RefSeq ORF:
 1632 bp

 Locus ID:
 4747

 UniProt ID:
 P07196

 Cytogenetics:
 8p21.2

Domains: filament, filament head

Protein Families: Druggable Genome, ES Cell Differentiation/IPS





Protein Pathways: Amyotrophic lateral sclerosis (ALS)

MW: 61.5 kDa

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, Oct 2008]