

Product datasheet for RC216128L2V

OriGene Technologies, Inc.

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NRG2 (NM_013982) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: NRG2 (NM_013982) Human Tagged ORF Clone Lentiviral Particle

Symbol: NRG2

Synonyms: DON1; HRG2; NTAK

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_013982 **ORF Size:** 2574 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC216128).

OTI Disclaimer:

Sequence:

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.

RefSeq: <u>NM 013982.1</u>

 RefSeq Size:
 3044 bp

 RefSeq ORF:
 2577 bp

 Locus ID:
 9542

 UniProt ID:
 014511

 Cytogenetics:
 5q31.2

Protein Families: Druggable Genome, Transmembrane

Protein Pathways: ErbB signaling pathway





ORIGENE

MW: 92.5 kDa

Gene Summary:

This gene encodes a novel member of the neuregulin family of growth and differentiation factors. Through interaction with the ERBB family of receptors, this protein induces the growth and differentiation of epithelial, neuronal, glial, and other types of cells. The gene consists of 12 exons and the genomic structure is similar to that of neuregulin 1, another member of the neuregulin family of ligands. The products of these genes mediate distinct biological processes by acting at different sites in tissues and eliciting different biological responses in cells. This gene is located close to the region for demyelinating Charcot-Marie-Tooth disease locus, but is not responsible for this disease. Alternative transcript variants encoding distinct isoforms have been described. [provided by RefSeq, May 2010]