

Product datasheet for RC227919L3V

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

NDE1 (NM_001143979) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: NDE1 (NM_001143979) Human Tagged ORF Clone Lentiviral Particle

Symbol: NDE1

Synonyms: HOM-TES-87; LIS4; MHAC; NDE; NUDE; NUDE1

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK

ACCN: NM_001143979

ORF Size: 1005 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC227919).

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.

RefSeq: <u>NM 001143979.1</u>, <u>NP 001137451.1</u>

 RefSeq Size:
 3936 bp

 RefSeq ORF:
 1008 bp

 Locus ID:
 54820

 UniProt ID:
 Q9NXR1

 Cytogenetics:
 16p13.11

 MW:
 37.7 kDa







Gene Summary:

This gene encodes a member of the nuclear distribution E (NudE) family of proteins. The encoded protein is localized at the centrosome and interacts with other centrosome components as part of a multiprotein complex that regulates dynein function. This protein plays an essential role in microtubule organization, mitosis and neuronal migration. Mutations in this gene cause lissencephaly 4, a disorder characterized by lissencephaly, severe brain atrophy, microcephaly, and severe cognitive disability. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Mar 2012]