

Product datasheet for RC202417L1V

OriGene Technologies, Inc.

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WNT7A (NM 004625) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: WNT7A (NM_004625) Human Tagged ORF Clone Lentiviral Particle

Symbol: Wnt-7a Synonyms: **Mammalian Cell**

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

Myc-DDK Tag: NM 004625 ACCN: **ORF Size:** 1047 bp

ORF Nucleotide

OTI Disclaimer:

Sequence:

The ORF insert of this clone is exactly the same as(RC202417).

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: NM 004625.3

RefSeq Size: 1732 bp RefSeq ORF: 1050 bp Locus ID: 7476 **UniProt ID:** <u>000755</u> Cytogenetics: 3p25.1 **Domains:**

Protein Families: Druggable Genome, Secreted Protein, Transmembrane





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Protein Pathways: Basal cell carcinoma, Hedgehog signaling pathway, Melanogenesis, Pathways in cancer, Wnt

signaling pathway

MW: 39 kDa

Gene Summary: This gene is a member of the WNT gene family, which consists of structurally related genes

that encode secreted signaling proteins. These proteins have been implicated in oncogenesis and in several developmental processes, including regulation of cell fate and patterning during embryogenesis. This gene is involved in the development of the anterior-posterior axis in the female reproductive tract, and also plays a critical role in uterine smooth muscle pattering and maintenance of adult uterine function. Mutations in this gene are associated

with Fuhrmann and Al-Awadi/Raas-Rothschild/Schinzel phocomelia syndromes. [provided by

RefSeq, Jul 2008]