

Product datasheet for RC206550L4

OriGene Technologies, Inc.

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Nkx2.5 (NKX2-5) (NM 004387) Human Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: Nkx2.5 (NKX2-5) (NM_004387) Human Tagged Lenti ORF Clone

Tag: mGFP Symbol: Nkx2.5

Synonyms: CHNG5; CSX; CSX1; HLHS2; NKX2.5; NKX2E; NKX4-1; VSD3

Mammalian Cell Puromycin

Selection:

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

E. coli Selection: Chloramphenicol (34 ug/mL)

ORF Nucleotide The ORF insert of this clone is exactly the same as(RC206550).

Sequence:

Restriction Sites: Sgfl-Mlul

Cloning Scheme:





^{*} The last codon before the Stop codon of the ORF

ACCN: NM_004387

ORF Size: 972 bp



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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.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

RefSeq: <u>NM 004387.2</u>

RefSeq Size: 1669 bp
RefSeq ORF: 975 bp
Locus ID: 1482

 UniProt ID:
 P52952

 Cytogenetics:
 5q35.1

Protein Families: Transcription Factors

MW: 34.9 kDa

Gene Summary: This gene encodes a homeobox-containing transcription factor. This transcription factor

functions in heart formation and development. Mutations in this gene cause atrial septal defect with atrioventricular conduction defect, and also tetralogy of Fallot, which are both

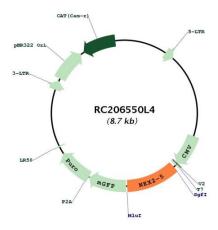
heart malformation diseases. Mutations in this gene can also cause congenital

hypothyroidism non-goitrous type 5, a non-autoimmune condition. Alternative splicing results

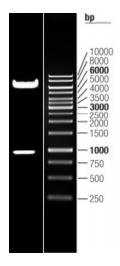
in multiple transcript variants. [provided by RefSeq, Oct 2009]



Product images:



Circular map for RC206550L4



Double digestion of RC206550L4 using Sgfl and Mul