

Product datasheet for MR223590L4

Dgcr6 (NM_010047) Mouse Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: Dgcr6 (NM_010047) Mouse Tagged Lenti ORF Clone

Tag: mGFP Symbol: Dgcr6

Mammalian Cell Puromycin

Selection:

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

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

ORF Nucleotide The ORF

Sequence:

The ORF insert of this clone is exactly the same as(MR223590).

Restriction Sites: Sgfl-Mlul

Cloning Scheme:





ACCN: NM_010047

ORF Size: 510 bp



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Dgcr6 (NM_010047) Mouse Tagged Lenti ORF Clone - MR223590L4

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 010047.3</u>, <u>NP 034177.1</u>

RefSeq Size: 1262 bp
RefSeq ORF: 513 bp
Locus ID: 13353
UniProt ID: 035347

Cytogenetics: 16 11.19 cM

Gene Summary: This gene encodes a protein that is similar to the gonadal protein in Drosophila (fruit fly). The

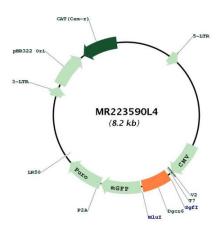
encoded protein is thought to play a role in migration of neural crest cells during development. Deletions in the human gene are associated with DiGeorge syndrome (or velocardiofacial syndrome) which has many clinical features including cardiac abnormalities, cleft palate, atypical facial features, hypocalcemia, hypoparathyroidism and defective

development or congenital absence of the thymus. Alternative splicing results in multiple

transcript variants. [provided by RefSeq, Jan 2014]



Product images:



Circular map for MR223590L4