

## Product datasheet for RC205805L4V

## OriGene Technologies, Inc.

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## QRX (RAX2) (NM\_032753) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** QRX (RAX2) (NM\_032753) Human Tagged ORF Clone Lentiviral Particle

Symbol: QRX

Synonyms: ARMD6; CORD11; QRX; RAXL1

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_032753

ORF Size: 552 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC205805).

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.

**RefSeg:** NM 032753.2

 RefSeq Size:
 2190 bp

 RefSeq ORF:
 555 bp

 Locus ID:
 84839

 UniProt ID:
 Q96IS3

 Cytogenetics:
 19p13.3

**Protein Families:** Transcription Factors

MW: 20.1 kDa







## **Gene Summary:**

This gene encodes a homeodomain-containing protein that plays a role in eye development. Mutation of this gene causes age-related macular degeneration type 6, an eye disorder resulting in accumulations of protein and lipid beneath the retinal pigment epithelium and within the Bruch's membrane. Defects in this gene can also cause cone-rod dystrophy type 11, a disease characterized by the initial degeneration of cone photoreceptor cells and resulting in loss of color vision and visual acuity, followed by the degeneration of rod photoreceptor cells, which progresses to night blindness and the loss of peripheral vision. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jan 2016]