

Product datasheet for RC213771L2V

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

ARX (NM 139058) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: ARX (NM_139058) Human Tagged ORF Clone Lentiviral Particle

Symbol:

CT121; EIEE1; ISSX; MRX29; MRX32; MRX33; MRX36; MRX38; MRX43; MRX54; MRX76; MRX87; Synonyms:

MRXS1; PRTS

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

mGFP Tag:

ACCN: NM 139058 ORF Size: 1686 bp

ORF Nucleotide

Sequence:

OTI Disclaimer:

The ORF insert of this clone is exactly the same as(RC213771).

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 139058.1

RefSeq Size: 2183 bp RefSeq ORF: 1689 bp Locus ID: 170302 **UniProt ID:** Q96QS3 Cytogenetics: Xp21.3

Domains: homeobox, OAR





ARX (NM_139058) Human Tagged ORF Clone Lentiviral Particle - RC213771L2V

Protein Families: Transcription Factors

MW: 58 kDa

Gene Summary: This gene is a homeobox-containing gene expressed during development. The expressed

protein contains two conserved domains, a C-peptide (or aristaless domain) and the prd-like class homeobox domain. It is a member of the group-II aristaless-related protein family whose members are expressed primarily in the central and/or peripheral nervous system. This gene is thought to be involved in CNS development. Expansion of a polyalanine tract and other mutations in this gene cause X-linked cognitive disability and epilepsy. [provided by

RefSeq, Jul 2016]