

Product datasheet for RC204899L2V

OriGene Technologies, Inc.

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Endothelin B Receptor (EDNRB) (NM 000115) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Endothelin B Receptor (EDNRB) (NM_000115) Human Tagged ORF Clone Lentiviral Particle

Symbol: Endothelin B Receptor

Synonyms: ABCDS; ET-B; ET-BR; ETB1; ETBR; ETRB; HSCR; HSCR2; WS4A

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_000115 **ORF Size:** 1326 bp

ORF Nucleotide

OTI Disclaimer:

1320 00

Sequence:

The ORF insert of this clone is exactly the same as(RC204899).

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 000115.1, NP 000106.1

 RefSeq Size:
 4296 bp

 RefSeq ORF:
 1329 bp

 Locus ID:
 1910

 UniProt ID:
 P24530

 Cytogenetics:
 13q22.3

Domains: 7tm 1

Protein Families: Druggable Genome, GPCR, Transmembrane





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Protein Pathways: Calcium signaling pathway, Melanogenesis, Neuroactive ligand-receptor interaction

MW: 49.6 kDa

Gene Summary: The protein encoded by this gene is a G protein-coupled receptor which activates a

phosphatidylinositol-calcium second messenger system. Its ligand, endothelin, consists of a family of three potent vasoactive peptides: ET1, ET2, and ET3. Studies suggest that the multigenic disorder, Hirschsprung disease type 2, is due to mutations in the endothelin receptor type B gene. Alternative splicing and the use of alternative promoters results in

multiple transcript variants. [provided by RefSeq, Oct 2016]