

Product datasheet for RC217313L1V

OriGene Technologies, Inc.

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ABCC11 (NM_032583) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: ABCC11 (NM_032583) Human Tagged ORF Clone Lentiviral Particle

Symbol: ABCC1²

Synonyms: EWWD; MRP8; WW

Mammalian Cell

Selection:

ACCN:

None

NM 032583

Vector: pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK

ORF Size: 4146 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC217313).

OTI Disclaimer:

Sequence:

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 032583.2

 RefSeq Size:
 4590 bp

 RefSeq ORF:
 4149 bp

 Locus ID:
 85320

 UniProt ID:
 Q96]66

 Cytogenetics:
 16q12.1

Protein Families: Druggable Genome, Transmembrane

Protein Pathways: ABC transporters





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MW: 154.1 kDa

Gene Summary:

The protein encoded by this gene is a member of the superfamily of ATP-binding cassette (ABC) transporters. ABC proteins transport various molecules across extra- and intra-cellular membranes. ABC genes are divided into seven distinct subfamilies (ABC1, MDR/TAP, MRP, ALD, OABP, GCN20, White). This ABC full transporter is a member of the MRP subfamily which is involved in multi-drug resistance. The product of this gene participates in physiological processes involving bile acids, conjugated steroids, and cyclic nucleotides. In addition, a SNP in this gene is responsible for determination of human earwax type. This gene and family member ABCC12 are determined to be derived by duplication and are both localized to chromosome 16q12.1. Multiple alternatively spliced transcript variants have been described for this gene. [provided by RefSeq, Jul 2008]