

Product datasheet for RC200949L1V

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

ABCB6 (NM_005689) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: ABCB6 (NM_005689) Human Tagged ORF Clone Lentiviral Particle

Symbol: ABCB6

Synonyms: ABC; LAN; MTABC3; PRP; umat

Mammalian Cell

Selection:

None

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

Tag: Myc-DDK
ACCN: NM 005689

ORF Size: 2526 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC200949).

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.

RefSeq: <u>NM 005689.1</u>

 RefSeq Size:
 3021 bp

 RefSeq ORF:
 2529 bp

 Locus ID:
 10058

 UniProt ID:
 Q9NP58

 Cytogenetics:
 2q35

Domains: ABC_membrane, ABC_tran, AAA

Protein Families: Druggable Genome, Transmembrane





ABCB6 (NM_005689) Human Tagged ORF Clone Lentiviral Particle - RC200949L1V

Protein Pathways: ABC transporters

MW: 93.9 kDa

Gene Summary: This gene encodes a member of the ATP-binding cassette (ABC) transporter superfamily. ABC

proteins transport various molecules across extra- and intra-cellular membranes. This protein is a member of the heavy metal importer subfamily and plays a role in porphyrin transport. This gene is the molecular basis of the Langereis (Lan) blood group antigen and mutations in this gene underlie familial pseudohyperkalemia and dyschromatosis universalis

hereditaria. [provided by RefSeq, Mar 2017]