

# Product datasheet for RC221445L4

# ATP8A2 (NM\_016529) Human Tagged Lenti ORF Clone

### **Product data:**

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Expression Plasmids
ATP8A2 (NM_016529) Human Tagged Lenti ORF Clone
mGFP
ATP8A2
ATP; ATPIB; CAMRQ4; IB; ML-1
Puromycin
pLenti-C-mGFP-P2A-Puro (PS100093)
Chloramphenicol (34 ug/mL)
The ORF insert of this clone is exactly the same as(RC221445).
Sgfl-Mlul
Cloning sites used for ORF Shuttling:
Sgf I         ORF         Mlu I            GCG ATC GC         ATG // NNÑ         ACG CGT



ACCN: ORF Size: NM\_016529 3564 bp



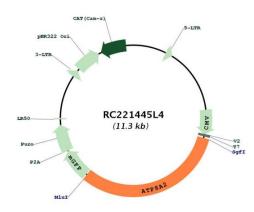
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	2 (NM_016529) Human Tagged Lenti ORF Clone – RC221445L4
OTI Disclaimer:	The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through 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. <u>More info</u>
OTI Annotation:	This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.
Components:	The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).
Reconstitution Method:	<ol> <li>Centrifuge at 5,000xg for 5min.</li> <li>Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li> <li>Close the tube and incubate for 10 minutes at room temperature.</li> <li>Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li> <li>Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.</li> </ol>
RefSeq:	<u>NM 016529.4</u>
RefSeq Size:	5006 bp
RefSeq ORF:	3567 bp
Locus ID:	51761
UniProt ID:	<u>Q9NTI2</u>
Cytogenetics:	13q12.13
Protein Families:	Transmembrane
MW:	133.4 kDa
Gene Summary:	The protein encoded by this gene is a member of the P4 ATPase family of proteins, which are thought to be involved in a process called lipid flipping, whereby phospholipids are translocated inwards from the exoplasmic leaflet to the cytosolic leaflet of the cell membrane, which aids in generating and maintaining asymmetry in membrane lipids. This protein is predicted to contain an E1 E2 ATPase, a haloacid dehalogenase-like hydrolase (HAD) domain, and multiple transmembrane domains. Associations between this protein and cell cycle control protein 50A are important for translocation of phosphatidylserine across membranes. Mutations in this gene have been associated with a syndrome (CAMRQ4) characterized by cerebellar ataxia and cognitive disabilities. In addition, a translocation breakpoint within this

gene was observed in an individual with neurological dysfunction. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Jul 2017]

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## **Product images:**



Circular map for RC221445L4

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