

Product datasheet for SC328756

LOXHD1 (NM 001173129) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: LOXHD1 (NM_001173129) Human Untagged Clone

Tag: Tag Free
Symbol: LOXHD1

Synonyms: DFNB77; LH2D1

Mammalian Cell

Selection:

None

Vector: pCMV6-XL5

E. coli Selection: Ampicillin (100 ug/mL)

Fully Sequenced ORF: >NCBI ORF sequence for NM_001173129, the custom clone sequence may differ by one or

more nucleotides

ATGACGGTGTGGACAGGGGATGTGGTTGGCGGGGGCACTGACTCCAACATCTTCATGACC GAGCGGGAGCAGAACGACACCTTCATCATGGAGATCCTAGACATTGCTCCATTCACCAAG ATGCGGATCCGGATTGATGGCCTGGGCAGTCGGCCGGAGTGGTTCCTGGAGAGGATCCTA CTGAAGAACATGAACACTGGAGACCTGACCATGTTCTACTATGGAGACTGGCTGTCCCAG CGGAAGGCCAAGAACCCTGGTGTGTGAAATGTGTGCCGTTATCGATGAGGAAGAAATG ATGGAGTGGACCTCCTACACCGTCGCAGTTAAGACCAGCGACATCCTGGGAGCAGGCACT GATGCCAACGTGTTCATCATCATCTTCGGGGAGAACGGGGATAGTGGGACACTGGCCCTG AAGCAGTCGGCAAACTGGAACAAGTTTGAGCGGAACAACACGGACACATTCAACTTCCCT GACATGCTGAGCTTGGGCCACCTCTGCAAGCTGAGGGTCTGGCACGACAACAAAGGGATA TTTCCTGGCTGGCATCTGAGCTATGTCGATGTGAAGGACAACTCCCGCGACGAGACCTTC CACTTCCAGTGTGACTGCTGGCTCTCCAAGAGTGAGGGTGACGGCAGACGGTCCGCGAC TTTGCCTGTGCCAACAACAAGATCTGTGATGAGCTGGAAGAGACCACCTACGAGATCGTC ATAGAAACGGGCAACGGAGGCGAAACCAGGGAGAACGTCTGGCTCATCCTGGAGGGCAGG AAGAACCGATCCAAAGAGTTTCTCATGGAAAATTCTTCTAGGCAGCGGGCCTTTAGGAAG GTGGGCCACCTTGCCAGGGAAGACCGGTTTATCCCCAAGAGAGAACTTGCCTGGCATGTC AAGACCATCACCATCACCGAGATGGAGTACGGCAATGTGTACTTCTTTAACTGTGACTGC CTCATCCCCCTCAAGAGGAAGAGGAAGTACTTCAAGGTATTCGAGGTTACCAAGACGACA GAGAGCTTTGCCAGCAAGGTCCAGAGCCTGGTGCCCGTCAAGTACGAAGTCATCGTGACA ACAGGCTATGAGCCAGGGGCAGGCACTGATGCCAACGTCTTCGTGACCATCTTTGGGGCC AACGGAGACACAGGCAAGCGGGAGCTGAAGCAGAAAATGCGCAACCTCTTCGAGCGGGGC AGCACAGACCGCTTCTTCCTGGAGACGCTGGAGCTGGGTGAGCTGCGCAAGTAG

Restriction Sites: Please inquire



OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com

LOXHD1 (NM_001173129) Human Untagged Clone - SC328756

ACCN: NM_001173129

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

OTI Annotation: This TrueClone is provided through our Custom Cloning Process that includes sub-cloning

into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.

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: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. 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.

RefSeq: <u>NM 001173129.1</u>, <u>NP 001166600.1</u>

 RefSeq Size:
 1853 bp

 RefSeq ORF:
 1374 bp

 Locus ID:
 125336

 UniProt ID:
 Q8IVV2

 Cytogenetics:
 18q21.1

Gene Summary: This gene encodes a highly conserved protein consisting entirely of PLAT

(polycystin/lipoxygenase/alpha-toxin) domains, thought to be involved in targeting proteins to

the plasma membrane. Studies in mice show that this gene is expressed in the

mechanosensory hair cells in the inner ear, and mutations in this gene lead to auditory defects, indicating that this gene is essential for normal hair cell function. Screening of human families segregating deafness identified a mutation in this gene which causes DFNB77, a progressive form of autosomal-recessive nonsyndromic hearing loss (ARNSHL). Alternatively

spliced transcript variants encoding different isoforms have been noted for this gene.

[provided by RefSeq, Mar 2010]

Transcript Variant: This variant (4) is missing many exons from the 5' end, and differs at the 5' and 3' ends compared to variant 1. This results in translation initiation from an in-frame

downstream AUG, and a shorter isoform (4) compared to isoform 1.