

Product datasheet for **RG231967**

CDHH (CDH13) (NM_001220492) Human Tagged ORF Clone

Product data:

Product Type: Expression Plasmids
Product Name: CDHH (CDH13) (NM_001220492) Human Tagged ORF Clone
Tag: TurboGFP
Symbol: CDH13
Synonyms: CDHH; P105
Mammalian Cell Selection: Neomycin
Vector: pCMV6-AC-GFP (PS100010)
E. coli Selection: Ampicillin (100 ug/mL)
ORF Nucleotide Sequence: >RG231967 representing NM_001220492
 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
 GCC**CGATCGCC**

ATGCAGCCGAGAACTCCGCTCGTTCGTGCGTTCTCCTGTCCCAGGTGCTGCTGCTAACATCTGCAGAAG
 ATTTGGACTGCACTCCTGGATTTACAGCAGAAAGTGTCCATATCAATCAGCCAGCTGAATTCATTGAGGA
 CCAGTCAATTCTAACTTGACCTTCAGTACTGTAAGGGAAACGACAAGCTACGCTATGAGGTCTCGAGC
 CCATACTCAAGGTGAACAGCGATGGCGGCTTAGTTGCTCTGAGAAACATAACTGCAGTGGGCAAAACTC
 TGTTTCGTCATGCACGGACCCCATGCGGAAGATATGGCAGAACTCGTGATTGTCGGGGGAAAGACAT
 CCAGGGCTCCTTGAGGATATATTTAAATTTGCAAGAACTTCTCCTGTCCAAGACAAAAGAGGTCCATT
 GTGGTATCTCCATTTAATTCCAGAGAATCAGAGACAGCCTTTCCCAAGAGATGTTGGCAAGATGAAGA
 TTTGGCAAGTTCTGTGCCTAGCACGATGGCTGACA

ACGCGTACGCGGCCGCTCGAG - GFP Tag - GTTTAA

Protein Sequence: >RG231967 representing NM_001220492
 Red=Cloning site Green=Tags(s)

MQPRTPLVLCVLLSQVLLLTSAEDLDCTPGFQQKVFHINQPAEFIEDQSILNLTFSACKGNDKLRVEVSS
 PYFKVNSDGGGLVALRNITAVGKTLFVHARTPHAEDMAELVIVGGKDIQGSQDIFKFARTSPVPRQKRSI
 VVSPILIPENRQPFPRDVGKMKIQQVLCARWLT

TRTRPLE - GFP Tag - V

Restriction Sites: SgfI-MluI



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OTI Disclaimer:	<p>Due to the inherent nature of this plasmid, standard methods to replicate additional amounts of DNA in E. coli are highly likely to result in mutations and/or rearrangements. Therefore, OriGene does not guarantee the capability to replicate this plasmid DNA. Additional amounts of DNA can be purchased from OriGene with batch-specific, full-sequence verification at a reduced cost. Please contact our customer care team at custsupport@origene.com or by calling 301.340.3188 option 3 for pricing and delivery.</p> <p>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</p>
OTI Annotation:	<p>This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.</p>
Components:	<p>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).</p>
Reconstitution Method:	<ol style="list-style-type: none">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:	NM_001220492.2
RefSeq Size:	1006 bp
RefSeq ORF:	528 bp
Locus ID:	1012
UniProt ID:	P55290
Cytogenetics:	16q23.3
Gene Summary:	<p>This gene encodes a member of the cadherin superfamily. The encoded protein is localized to the surface of the cell membrane and is anchored by a GPI moiety, rather than by a transmembrane domain. The protein lacks the cytoplasmic domain characteristic of other cadherins, and so is not thought to be a cell-cell adhesion glycoprotein. This protein acts as a negative regulator of axon growth during neural differentiation. It also protects vascular endothelial cells from apoptosis due to oxidative stress, and is associated with resistance to atherosclerosis. The gene is hypermethylated in many types of cancer. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, May 2011]</p>