

## Product datasheet for **RG204202**

### Claudin 14 (CLDN14) (NM\_012130) Human Tagged ORF Clone

#### Product data:

Product Type:	Expression Plasmids
Product Name:	Claudin 14 (CLDN14) (NM_012130) Human Tagged ORF Clone
Tag:	TurboGFP
Symbol:	Claudin 14
Synonyms:	DFNB29
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-AC-GFP (PS100010)
E. coli Selection:	Ampicillin (100 ug/mL)
ORF Nucleotide Sequence:	>RG204202 representing NM_012130 Red=Cloning site Blue=ORF Green=Tags(s)

TTTTGTAATACGACTCACTATAGGGCGGCCGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC  
GCC**CGATCGCC**

ATGGCCAGCACGGCCGTGCAGCTTCTGGGCTTCTGCTCAGCTTCTGGGCATGGTGGGCACGTTGATCA  
CCACCATCCTGCCGACTGGCGGAGGACAGCGCACGTGGGCACCAACATCCTCACGGCCGTCTACCTACCT  
GAAAGGGCTCTGGATGGAGTGTGTGGCACAGCACAGGCATCTACCAGTGCCAGATCTACCGATCCCTG  
CTGGCGCTGCCCAAGACCTCCAGGCTGCCCGGCCCTCATGGTCATCTCCTGCCTGCTCTCGGGCATAG  
CCTGCGCCTGCGCCGTATCGGGATGAAGTGCACGCGCTGCGCCAAGGGCACACCCGCCAAGACCACCTT  
TGCCATCCTCGGCGGCACCCTTTCATCCTGGCCGGCCTCCTGTGCATGGTGGCCGTCTCCTGGACCACC  
AACGACGTGGTGCAGAACTTCTACAACCCGCTGCTGCCAGCGGCATGAAGTTTGAGATTGGCCAGGCC  
TGTACCTGGGCTTCATCTCCTCGTCCCTCTCGCTCATTGGTGGCACCCCTGCTTTGCCTGTCTGCCAGGA  
CGAGGCACCCTACAGGCCCTACCAGGCCCGCCAGGGCCACCACGACCACTGCAAAACCCGCACCTGCC  
TACCAGCCACCAGCTGCCTACAAAGACAATCGGGCCCCCTCAGTGACCTCGGCCACGCACAGCGGGTACA  
GGCTGAACGACTACGTG

**ACGCGT**ACGCGGCCGCTCGAG - GFP Tag - GTTTAA



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**Protein Sequence:** >RG204202 representing NM\_012130  
 Red=Cloning site Green=Tags(s)

MASTAVQLLGFLLSFLGMVGLITTTILPHWRRTAHVGTNILTAVSYLKGLWMECVHSTGIYQCQIYRSL  
 LALPQDLQAARALMVISCLLSGIACACAVIGMKTRCAKGTAKTTFFAILGGTLFILAGLLCMVAVSWTT  
 NDVVQNFYNPLLPSGMKFEIGQALYLGFISSSLIGGTLLCLSCQDEAPYRPYQAPPRATTTTANTAPA  
 YQPPAAAYKDNRAPSVTSATHSGYRLNDYV

TRTRPLE - GFP Tag - V

**Restriction Sites:** SgfI-MluI

**Cloning Scheme:**



**ACCN:** NM\_012130

**ORF Size:** 717 bp

**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.

**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:** [NM\\_012130.4](#)

**RefSeq Size:** 1233 bp

**RefSeq ORF:** 720 bp

**Locus ID:** 23562

**UniProt ID:** [O95500](#)

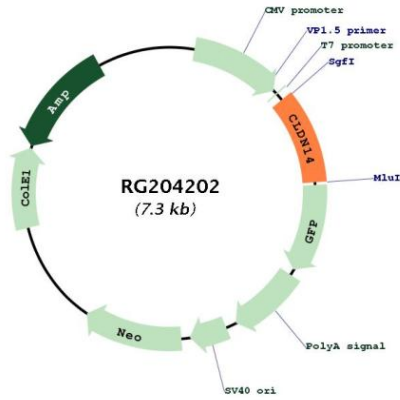
**Cytogenetics:** 21q22.13

**Protein Families:** Druggable Genome, Transmembrane

**Protein Pathways:** Cell adhesion molecules (CAMs), Leukocyte transendothelial migration, Tight junction

**Gene Summary:** Tight junctions represent one mode of cell-to-cell adhesion in epithelial or endothelial cell sheets, forming continuous seals around cells and serving as a physical barrier to prevent solutes and water from passing freely through the paracellular space. These junctions are comprised of sets of continuous networking strands in the outwardly facing cytoplasmic leaflet, with complementary grooves in the inwardly facing extracytoplasmic leaflet. The protein encoded by this gene, a member of the claudin family, is an integral membrane protein and a component of tight junction strands. The encoded protein also binds specifically to the WW domain of Yes-associated protein. Defects in this gene are the cause of an autosomal recessive form of nonsyndromic sensorineural deafness. It is also reported that four synonymous variants in this gene are associated with kidney stones and reduced bone mineral density. Several transcript variants encoding the same protein have been found for this gene. [provided by RefSeq, Jun 2010]

Product images:



Circular map for RG204202