

Full-Length

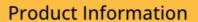
CLDN14 (Human) Recombinant Protein (P01)

Catalog # H00023562-P01 Size 25 ug, 10 ug

Applications



Specification	
Product Description	Human CLDN14 full-length ORF (NP_036262.1, 1 a.a 239 a.a.) recombinant protein with GST-tag at N-terminal.
Sequence	MASTAVQLLGFLLSFLGMVGTLITTILPHWRRTAHVGTNILTAVSYLKGLWMECVWHSTGIYQCQIY RSLLALPQDLQAARALMVISCLLSGIACACAVIGMKCTRCAKGTPAKTTFAILGGTLFILAGLLCMV AVSWTTNDVVQNFYNPLLPSGMKFEIGQALYLGFISSSLSLIGGTLLCLSCQDEAPYRPYQAPPRA TTTTANTAPAYQPPAAYKDNRAPSVTSATHSGYRLNDYV
Host	Wheat Germ (in vitro)
Theoretical MW (kDa)	52.1
Interspecies Antigen Sequence	Mouse (93); Rat (93)
Preparation Method	in vitro wheat germ expression system
Purification	Glutathione Sepharose 4 Fast Flow
Quality Control Testing	12.5% SDS-PAGE Stained with Coomassie Blue.
Storage Buffer	50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.





Note

Best use within three months from the date of receipt of this protein.

Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

Gene Info — CLDN14	
Entrez GeneID	23562
GeneBank Accession#	NM_012130.2
Protein Accession#	NP_036262.1
Gene Name	CLDN14
Gene Alias	DFNB29
Gene Description	claudin 14
Omim ID	<u>605608</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	Tight junctions represent one mode of cell-to-cell adhesion in epithelial or endothelial cell sheets, f orming continuous seals around cells and serving as a physical barrier to prevent solutes and wat er 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. Several transcript variants encoding the same protein have been found for this gene. [provided by RefSeq
Other Designations	OTTHUMP00000109045 OTTHUMP00000109046 OTTHUMP00000109047 OTTHUMP00000109048 OTTHUMP00000109049



Pathway

- Cell adhesion molecules (CAMs)
- Leukocyte transendothelial migration
- Tight junction

Disease

- Genetic Predisposition to Disease
- Hearing Loss
- Kidney Calculi
- Tobacco Use Disorder