

Proteoliposomes

Full-Length

OPN1MW (Human) Recombinant Protein

Catalog # H00002652-G01

Size 2 ug

Specification

Product Description	Human OPN1MW full-length ORF (AAI56777.1) recombinant protein without tag. This product is belong to Proteoliposome (PL).
Sequence	MAQQWSLQRLAGRHPQDSYEDSTQSSIFTYTNSNSTRGPFEGPNYHIAPRWVYHLTSVWMIFVVI ASVFTNGLVLAATMKFKLRHPLNWILVNLAVADLAETVIASISVVNQVYGYFVLGHMPCVLEGYT VSLCGITGLWSLAISWERWMVCKPFGNVRFDKLAIVGIAFSWMAAVWTAPPIFGWSRYWPH GLKTSCGPDVFSGSSYPGVQSYMMLMVTCITPLSIMLCYLQVWLAIKRAVAKQQKESESTQKAEK EVTRMVVVMVLAFCFCWGPYAFFACFAAANPGYPFHPLMAALPAFFAKSATIYNPVIYVFMNRQF RNCILQLFGKKVDDGSELSSASKTEVSSVSSVSPA
Host	Wheat Germ (in vitro)
Theoretical MW (kDa)	40.04
Form	Liquid
Preparation Method	in vitro wheat germ expression system with proprietary liposome technology
Purification	None
Recommend Usage	Heating may cause protein aggregation. Please do not heat this product before electrophoresis.
Storage Buffer	25 mM Tris-HCl of pH8.0 containing 2% glycerol.
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

- Antibody Production

Gene Info — OPN1MW

Entrez GeneID [2652](#)

GeneBank Accession# [BC156776.1](#)

Protein Accession# [AA156777.1](#)

Gene Name OPN1MW

Gene Alias CBBM, CBD, GCP, MGC176615, OPN1MW1

Gene Description opsin 1 (cone pigments), medium-wave-sensitive

Omim ID [303700](#) [303800](#)

Gene Ontology [Hyperlink](#)

Gene Summary

This gene encodes for a light absorbing visual pigment of the opsin gene family. The encoded protein is called green cone photopigment or medium-wavelength sensitive opsin. Opsins are G-protein coupled receptors with seven transmembrane domains, an N-terminal extracellular domain, and a C-terminal cytoplasmic domain. The long-wavelength opsin gene and multiple copies of the medium-wavelength opsin gene are tandemly arrayed on the X chromosome and frequent unequal recombination and gene conversion may occur between these sequences. X chromosomes may have fusions of the medium- and long-wavelength opsin genes or may have more than one copy of these genes. Defects in this gene are the cause of deutanopic colorblindness. [provided by RefSeq]

Other Designations

OTTHUMP00000024237|color blindness, deutan|green cone photoreceptor pigment|green cone pigment|opsin 1 (cone pigments), medium-wave-sensitive (color blindness, deutan)|photopigment apoprotein