

DNAxPAb



# **OPN1LW DNAxPab**

Catalog # H00005956-W01P Size 200 ug

Specification	
Product Description	Rabbit polyclonal antibody raised against a partial-length human OPN1LW DNA using DNAx™ Immu ne technology.
Technology	<u>DNAx™ Immune</u>
Immunogen	Extracellular membrane domain (ECD) human DNA
Host	Rabbit
Reactivity	Human
Purification	Protein A
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

#### Applications

- Western Blot (Transfected lysate)
  <u>Protocol Download</u>
- Immunofluorescence (Transfected cell)
- Flow Cytometry (Transfected cell)

### Gene Info — OPN1LW

😵 Abnova

## **Product Information**

Entrez GenelD	<u>5956</u>
GeneBank Accession#	BC156643.1
Protein Accession#	<u>AAI56644.1</u>
Gene Name	OPN1LW
Gene Alias	CBBM, CBP, RCP
Gene Description	opsin 1 (cone pigments), long-wave-sensitive
Omim ID	<u>303700</u> <u>303900</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes for a light absorbing visual pigment of the opsin gene family. The encoded pro tein is called red cone photopigment or long-wavelength sensitive opsin. Opsins are G-protein co upled receptors with seven transmembrane domains, an N-terminal extracellular domain, and a C -terminal cytoplasmic domain. This gene and the medium-wavelength opsin gene are tandemly ar rayed 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-wavelengt h opsin genes or may have more than one copy of these genes. Defects in this gene are the caus e of partial, protanopic colorblindness. [provided by RefSeq
Other Designations	OTTHUMP00000032193 red cone photoreceptor pigment red-sensitive opsin

#### Disease

- Genetic Predisposition to Disease
- <u>Retinal Diseases</u>
- Sarcoidosis