

Product Information

MemDX™ Membrane Protein Human CLN6 (CLN6 transmembrane ER protein) for Antibody

Discovery

Cat. No.: **MP0507J**

This product is for research use only and is not intended for diagnostic use.

This product is a 35.7 kDa Human CLN6 membrane protein expressed in HEK293T. The protein is for research use only and is not approved for use in humans or in clinical diagnosis.

Product Specifications

Host Species

Human

Target Protein

CLN6

Protein Length

Full-length

Protein Class

Transmembrane

Molecular Weight

35.7 kDa

TMD

7

Sequence

MEATRRRQHLGATGGPGAQLGASFLQARHGVSADAEARTAPFHLDLWFYFTLQNWVLDGFRPIAMLVFP
LEWFPLNKPSVGDYFHMAYNVITPFLLLKLIERSPRTLPRSITYVSIIFIMGASIHVGDVSNHRLIFS
GYQHHLVSVRENPIIKNLKPETLIDSFELLYYYDEYLGHCMWYIPFFLILFMYFSGCFTASKAESLIPGPA
LLLVAAPSGLYYWYLVTEGQIFILFIFTFFAMLALVLHQKRKRLFLDSNGLFLFSSFALTLLLVALWVAWL
WNDPVLRRKKYPGVIYVPEPWAFYTLHVSSRH

Product Description

Expression Systems

HEK293T

Tag

C-Myc/DDK

Form

Liquid

Purification

Anti-DDK affinity column followed by conventional chromatography steps

Purity

> 80% as determined by SDS-PAGE and Coomassie blue staining

Buffer

25 mM Tris.HCl, pH 7.3, 100 mM glycine, 10% glycerol

Storage

Store at +4°C for up to one week or several months at -80°C

Target**Target Protein**

CLN6

Full Name

CLN6 transmembrane ER protein

Introduction

This gene is one of eight which have been associated with neuronal ceroid lipofuscinoses (NCL). Also referred to as Batten disease, NCL comprises a class of autosomal recessive, neurodegenerative disorders affecting children. The genes responsible likely encode proteins involved in the degradation of post-translationally modified proteins in lysosomes. The primary defect in NCL disorders is thought to be associated with lysosomal storage function.

Alternative Names

CLN4A; HsT18960; nclf

Gene ID

[54982](#)

UniProt ID

[Q9NWW5](#)