

Product Information

MemDX™ Membrane Protein Human WFS1 (Wolframin ER transmembrane glycoprotein) for Antibody Discovery

Cat. No.: **MP0656J**

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

This product is a 100.1 kDa Human WFS1 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

WFS1

Protein Length

Full-length

Protein Class

Druggable Genome, Transmembrane

Molecular Weight

100.1 kDa

TMD

11

Sequence

MDSNTAPLGPSCPPPPAPQPQARSRLNATASLEQERSERPRAPGPQAGPGPGVRDAAAPAEPQAQHTRS
RERADGTGPTKGDMEIPFEEVLERAKAGDPKAQTEVGKHYLQLAGDTDEELNSCTAVDWLVLAQKGRRE
AVKLLRRCLADRRGITSENEREVRQLSSETDLERAVRKAALVMYWKLNPKKKKQVAVAELENVGVNEH
DGGAQPGVPKSLQKQRRMLERLVSSSESKNYIALDDFVEITKKYAKGVIPSSLFLQDDEDDDELAGKSP
DLPLRLKVVKYPLHAIMEIKEYLIDMASRAGMHWLSTIPTHINALIFFFIVSNLTIDFFAFFIPLVIF
YLSFISMVICTLKVFQDSKAWENFRTLTDLLRFEPNLDVEQAEVNFNGWNHLEPYAHFLLSVFFVIFSFP
IASKDCIPCSELAVITGFFTTSYLSLSTHAEPYTRRALATEVTAGLLSLLPSMPLNWPYLVKVLGQTFIT
VPVGHVVLNVSVPCLLYVYLLYLFFRMAQLRNFKGTICYLVPYLVCFMWCELSVVILLESTGLGLLRAS
IGYFLFLFALPILVAGLALVGVLFARWFTSLELT KIAVTAVCSVPLLLRWWTKASFVVGMMVKSLTRS
SMVKLILVWLTAVLFCWFYVYRSEGMKVYNSTLTWQQYGALCGPRAWKETNMARTQILCSHLEGHRVTW
TGRFKYVRVTDIDNSAESAINMLPFFIGDWMRCLYGEAYPACSPGNTSTAEELCRLKLLAKHPCHIKKF
DRYKFEITVGMPSFGADGSRSDREDDVTKDIVLRASSEFKSVLLSLRQGSLEIFSTILEGRLGSKWPVF
ELKAISCLNCMAQLSPTRRHVKIEHDWRSTVHGAVKFAFDFFFFPFLSAA

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**

WFS1

Full Name

Wolframin ER transmembrane glycoprotein

Introduction

This gene encodes a transmembrane protein, which is located primarily in the endoplasmic reticulum and ubiquitously expressed with highest levels in brain, pancreas, heart, and insulinoma beta-cell lines. Mutations in this gene are associated with Wolfram syndrome, also called DIDMOAD (Diabetes Insipidus, Diabetes Mellitus, Optic Atrophy, and Deafness), an autosomal recessive disorder. The disease affects the brain and central nervous system. Mutations in this gene can also cause autosomal dominant deafness 6 (DFNA6), also known as DFNA14 or DFNA38. Alternatively spliced transcript variants have been found for this gene.

Alternative Names

WFS; WFRS; WFSL; CTRCT41

Gene ID

[7466](#)

UniProt ID

[Q76024](#)