

Product Information

MemDX™ Membrane Protein Human FA2H (Fatty acid 2-hydroxylase) for Antibody

Discovery

Cat. No.: **MP0493J**

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

This product is a 42.6 kDa Human FA2H 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

FA2H

Protein Length

Full-length

Protein Class

Transmembrane

Molecular Weight

42.6 kDa

TMD

4

Sequence

MAPAPPPAASFSPSEVQRRRLAAGACWVRRGARLYDLSSFVRHHPGGEQLLRARAGQDISADLDGPPHRHS
ANARRWLEQYYVGELRGEQQGSMENEPVALEETQKTDPA MEPRFKVVDWDKDLVDWRKPLLWQVGH LGEK
YDEWVHQPVTRPIRLFHSDLIEGLSKTVWYSVP IIVVPLVLYLSWSYYRTFAQGNVRLFTSFTTEYTVAV
PKSMFPGFLFMLGTFLWLSLIEYLIHRFLFHMKPPSDSYYLIMLHFVMH GQH HKAPFDGSRLVFPPVPASLV
IGVFYLCMQLILPEAVGGTVFAGGLLGYVLYDMTHYYLHFGSPHKGSYLYSLKAHHVKHHFAHQKSGFGI
STKLWDYCFHTLTPEKPHLKTQ

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**

FA2H

Full Name

Fatty acid 2-hydroxylase

Introduction

This gene encodes a protein that catalyzes the synthesis of 2-hydroxysphingolipids, a subset of sphingolipids that contain 2-hydroxy fatty acids. Sphingolipids play roles in many cellular processes and their structural diversity arises from modification of the hydrophobic ceramide moiety, such as by 2-hydroxylation of the N-acyl chain, and the existence of many different head groups. Mutations in this gene have been associated with leukodystrophy dysmyelinating with spastic paraparesis with or without dystonia.

Alternative Names

FAAH; FAH1; SCS7; SPG35; FAXDC1

Gene ID

[79152](#)

UniProt ID

[Q7L5A8](#)