Recombinant Human FA2H, His-tagged

Cat. No. FA2H-12628H **Lot. No.** (See product label)

SPECIFICATION

Product Overview	Recombinant Human FA2H protein, fused to His-tag, was expressed in E.coli and purified by Ni-sepharose.
Species	Human
Source	E.coli
ProteinLength	1-164a.a.
Description	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.
Storage	The protein is stored in PBS buffer at -20℃. Avoid repeated freezing and thawing cycles.
Storage Buffer	1M PBS (58mM Na2HPO4,17mM NaH2PO4, 68mM NaCl, pH8.) added with 300mM Imidazole and 0.7% Sarcosyl, 15%glycerol.

GENE INFORMATION

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Gene Name	FA2H fatty acid 2-hydroxylase [Homo sapiens]
Official Symbol	FA2H
Synonyms	FA2H; fatty acid 2-hydroxylase; fatty acid hydroxylase domain containing 1 , FAXDC1, spastic paraplegia 35 (autosomal recessive) , SPG35; FAAH; fatty acid hydroxylase; FLJ25287; fatty acid alpha-hydroxylase; fatty acid hydroxylase domain containing 1; spastic paraplegia 35 (autosomal recessive); FAH1; SCS7; SPG35; FAXDC1;
Gene ID	79152
mRNA Refseq	NM_024306
Protein Refseq	NP_077282
MIM	611026
UniProt ID	Q7L5A8
Chromosome Location	16q23
Function	heme binding; metal ion binding; oxidoreductase activity;

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