

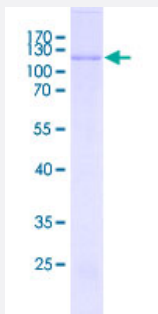
Full-Length

MFN2 (Human) Recombinant Protein (P01)

Catalog # H00009927-P01

Size 10 ug, 25 ug

Applications



Specification

Product Description

Human MFN2 full-length ORF (NP_055689.1, 1 a.a. - 757 a.a.) recombinant protein with GST tag at N-terminal.

Sequence

MSLLFSRCNSIVTKKNKRHMAEVNASPLKHFVTAKKKINGIFEQLGAYIQESATFLEDTYRNAELD
PVTTEEQVLDVKGYLSKVRGISEVLARRHMKVAFFGRTSNGKSTVINAMLWDKVLPSGIGHTTNC
FLRVEGTDGHEAFLATEGSEEKRSKATVNQLAHLHQDKQLHAGSLVSVMWPNSKCPLLKDDL
VLMDSPGIDVTTELD SWIDKFCLDADVFVLVANSESTLMQTEKHFFHKVSERLSRPNFILNNRWD
ASASEPEYMEEVRRQHMERCTSFLVDELGVVDRSQAGDRIFVSAKEVLNARIQKAQGMPEGG
GALAEGFQVRMFEFQNFERRFEECISQSAVKTKFEQHTVRAKQIAEAVRLIMDSLHMAAREQQVY
CEEMREERQDRLKFIDKQLELLAQDYKLRIQITEEVERQVSTAMAEIIRLSVLVDDYQMDFHPS
PVVLKVYKNELHRHIEEGLGRNMSDRCASTATNSLQTMQQDMIDGLKPLLPVSVRSQIDMLVPRQC
FSLNYDLNCDKLCADFQEDIEFHFSLGWTMLVNRFLGPKNSRRALMGYNDQVQRPIPLTPANPS
MPPLPQGSLTQEEFMVSMVTGLASLTSRTSMGILVGGVWVKAVGWRLIALSFGLYGLLYYERL
TWTTKAKERAFKRQFVEHASEKLQLVISYTGSNCSHQVQQELSGTFAHLCQQVDVTRENLEQEI
AMNKKIEVLDSLQSKAKLLRNKAGWLDSELNMFTHQYLQPSR

Host

Wheat Germ (in vitro)

Theoretical MW (kDa)

112.8

Interspecies Antigen Sequence

Mouse (95); Rat (96)

Preparation Method

[in vitro wheat germ expression system](#)

Purification	Glutathione Sepharose 4 Fast Flow
Quality Control Testing	12.5% SDS-PAGE Stained with Coomassie Blue
Storage Buffer	50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
Note	Best use within three months from the date of receipt of this protein.

Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

Gene Info — MFN2

Entrez GeneID	9927
GeneBank Accession#	NM_014874.2
Protein Accession#	NP_055689.1
Gene Name	MFN2
Gene Alias	CMT2A, CMT2A2, CPRP1, HSG, KIAA0214, MARF
Gene Description	mitofusin 2
Omim ID	601152 608507 609260
Gene Ontology	Hyperlink

Gene Summary	This gene encodes a mitochondrial membrane protein that participates in mitochondrial fusion and contributes to the maintenance and operation of the mitochondrial network. This protein is involved in the regulation of vascular smooth muscle cell proliferation, and it may play a role in the pathophysiology of obesity. Mutations in this gene cause Charcot-Marie-Tooth disease type 2A2, and hereditary motor and sensory neuropathy VI, which are both disorders of the peripheral nervous system. Defects in this gene have also been associated with early-onset stroke. Two transcript variants encoding the same protein have been identified. [provided by RefSeq]
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Other Designations

OTTHUMP00000002509|hyperplasia suppressor|mitochondrial assembly regulatory factor|mitofusin-2|transmembrane GTPase MFN2

Disease

- [Charcot-Marie-Tooth Disease](#)
- [Genetic Predisposition to Disease](#)
- [Glaucoma](#)
- [Hereditary Sensory and Motor Neuropathy](#)