

MFN2 FISH Probe

Catalog # FA0012 Size 200 uL

Specification	
Product Description	Made to order FISH probes for identification of gene amplification using Fluorescent In Situ Hybridiz ation Technique. (Technology).
Origin	Human
Source	Genomic DNA
Reactivity	Human
Notice	We strongly recommend the customer to use FFPE FISH PreTreatment Kit 1 (Catalog #: <u>KA2375</u> or <u>KA2691</u>) for the pretreatment of Formalin-Fixed Paraffin-Embedded (FFPE) tissue sections.
Regulation Status	For research use only (RUO)
Supplied Product	DAPI Counterstain (1500 ng/mL) 250 uL
Storage Instruction	Store at 4°C in the dark.

Applications

• Fluorescent In Situ Hybridization (Cell)

Protocol Download

Gene Info — MFN2	
Entrez GenelD	<u>9927</u>
Gene Name	MFN2
Gene Alias	CMT2A, CMT2A2, CPRP1, HSG, KIAA0214, MARF
Gene Description	mitofusin 2

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Product Information

Omim ID	<u>601152 608507 609260</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a mitochondrial membrane protein that participates in mitochondrial fusion an d contributes to the maintenance and operation of the mitochondrial network. This protein is involv ed in the regulation of vascular smooth muscle cell proliferation, and it may play a role in the patho physiology of obesity. Mutations in this gene cause Charcot-Marie-Tooth disease type 2A2, and h ereditary motor and sensory neuropathy VI, which are both disorders of the peripheral nervous sys tem. Defects in this gene have also been associated with early-onset stroke. Two transcript varian ts encoding the same protein have been identified. [provided by RefSeq
Other Designations	OTTHUMP0000002509 hyperplasia suppressor mitochondrial assembly regulatory factor mitofu sin-2 transmembrane GTPase MFN2

Disease

- <u>Charcot-Marie-Tooth Disease</u>
- Genetic Predisposition to Disease
- Glaucoma
- Hereditary Sensory and Motor Neuropathy