

## Mitofusin 2 Polyclonal Antibody

Catalog No. E-AB-64745

**Note:** Centrifuge before opening to ensure complete recovery of vial contents.

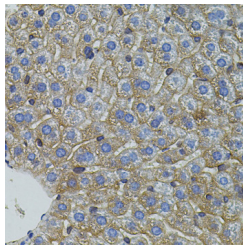
### Description

|                     |  |
|---------------------|--|
| <b>Reactivity</b>   | Human,Mouse,Rat  |
| <b>Immunogen</b>    | Recombinant fusion protein of mouse Mitofusin 2 (NP_573464.2). |
| <b>Host</b>         | Rabbit   |
| <b>Isotype</b>      | IgG  |
| <b>Purification</b> | Affinity purification  |
| <b>Conjugation</b>  | Unconjugated   |
| <b>Buffer</b>       | PBS with 0.02% sodium azide, 50% glycerol, pH7.3.              |

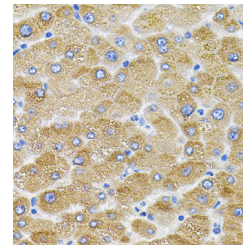
### Applications Recommended Dilution

IHC 1:50-1:200

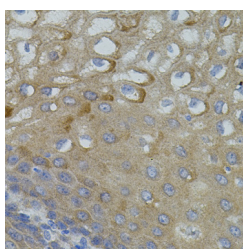
### Data



Immunohistochemistry of paraffin-embedded Mouse liver using Mitofusin 2 Polyclonal Antibody



Immunohistochemistry of paraffin-embedded Human liver damage using Mitofusin 2 Polyclonal Antibody



Immunohistochemistry of paraffin-embedded Human esophagus using Mitofusin 2 Polyclonal Antibody

### Preparation & Storage

**Storage** Store at -20°C. Avoid freeze / thaw cycles.

### Background

This gene encodes a mitochondrial membrane protein that participates in mitochondrial fusion and contributes to the

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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.