MT-ND4L gene
mitochondrially encoded NADH:ubiquinone oxidoreductase core subunit 4L

Normal Function

The *MT-ND4L* gene provides instructions for making a protein called NADH dehydrogenase 4L. This protein is part of a large enzyme complex known as complex I, which is active in mitochondria. Mitochondria are structures within cells that convert the energy from food into a form that cells can use. These cellular structures produce energy through a process called oxidative phosphorylation, which uses oxygen and simple sugars to create adenosine triphosphate (ATP), the cell's main energy source.

Complex I is one of several enzyme complexes necessary for oxidative phosphorylation. Within mitochondria, these complexes are embedded in a tightly folded, specialized membrane called the inner mitochondrial membrane. During oxidative phosphorylation, mitochondrial enzyme complexes carry out chemical reactions that drive the production of ATP. Specifically, they create an unequal electrical charge on either side of the inner mitochondrial membrane through a step-by-step transfer of negatively charged particles called electrons. This difference in electrical charge provides the energy for ATP production.

Complex I is responsible for the first step in the electron transport process, the transfer of electrons from a molecule called NADH to another molecule called ubiquinone. Electrons are then passed from ubiquinone through several other enzyme complexes to provide energy for the generation of ATP.

Health Conditions Related to Genetic Changes

**Leber hereditary optic neuropathy**

A mutation in the *MT-ND4L* gene has been identified in several families with Leber hereditary optic neuropathy. This mutation, which can be written as T10663C or Val65Ala, changes a single protein building block (amino acid) in the NADH dehydrogenase 4L protein. Specifically, it replaces the amino acid valine with the amino acid alanine at protein position 65.

Researchers have not determined how a mutation in the *MT-ND4L* gene can lead to the vision loss characteristic of Leber hereditary optic neuropathy. This genetic change appears to disrupt the normal activity of complex I in the mitochondrial inner membrane, which may affect the production of ATP. It remains unclear, however, why the effects of this mutation are limited to the nerve that relays visual information from the eye to the brain (the optic nerve). Additional genetic and environmental factors probably contribute to the features of Leber hereditary optic neuropathy.
Mitochondrial complex I deficiency

Chromosomal Location

Other Names for This Gene

- Complex I, subunit ND4L
- mitochondrially encoded NADH 4L
- mitochondrially encoded NADH dehydrogenase 4L
- MTND4L
- NADH dehydrogenase 4L
NADH dehydrogenase subunit 4L
NADH-ubiquinone oxidoreductase chain 4L
NADH-ubiquinone oxidoreductase, subunit ND4L
NADH4L
ND4L
NU4LM_HUMAN

Additional Information & Resources

Educational Resources
- Mayo Clinic: North American Mitochondrial Disease Consortium Patient Registry and Biorepository (NAMDC)
  https://www.mayo.edu/research/clinical-trials/cls-20409244
- Oxidative Phosphorylation (Biochemistry, Fifth Edition, 2002)
  https://www.ncbi.nlm.nih.gov/books/NBK21208/
- The Neuromuscular Disease Center at Washington University: Complex I
  https://neuromuscular.wustl.edu/pathol DIAGRAMS/mito.htm#complexI

Clinical Information from GeneReviews
- Leber Hereditary Optic Neuropathy
  https://www.ncbi.nlm.nih.gov/books/NBK1174
- Mitochondrial Disorders Overview
  https://www.ncbi.nlm.nih.gov/books/NBK1224

Scientific Articles on PubMed
- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28MT-ND4L%5BTIAB%5D%29+OR+%28mitochondrially+encoded+NADH+4L%5BTIAB%5D%29%29+OR+%28%28MTND4L%5BTIAB%5D%29+OR+%28NADH+dehydrogenase+subunit+4L%5BTIAB%5D%29+OR+%28Complex+I,+subunit+ND4L%5BTIAB%5D%29+OR+%28NADH+dehydrogenase+4L%5BTIAB%5D%29+OR+%28NADH-ubiquinone+oxidoreductase+chain+4L%5BTIAB%5D%29+OR+%28NADH-ubiquinone+oxidoreductase,+subunit+ND4L%5BTIAB%5D%29+OR+%28NADH4L%5BTIAB%5D%29+OR+%28ND4L%5BTIAB%5D%29+AND +%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bl%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM
- COMPLEX I, SUBUNIT ND4L
  http://omim.org/entry/516004

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Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology
  http://atlasgeneticsoncology.org/Genes/GC_ND4L.html

- ClinVar

- HGNC Gene Symbol Report

- Mitomap: Leber Hereditary Optic Neuropathy Disease Mutation Database
  https://www.mitomap.org/MITOMAP/MutationsLHON

- Monarch Initiative
  https://monarchinitiative.org/gene/NCBIGene:4539

- NCBI Gene

- UniProt
  https://www.uniprot.org/uniprot/P03901

Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/11935318

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15972314
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2564640/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/14617834

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20301353

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