OPA1 gene

OPA1, mitochondrial dynamin like GTPase

Normal Function

The *OPA1* gene provides instructions for making a protein that is found in cells and tissues throughout the body. The OPA1 protein is active in the inner membrane of cell structures called mitochondria, which are the energy-producing centers in cells. Mitochondria are dynamic structures that change shape through processes called fission (splitting into smaller pieces) and fusion (combining pieces). Changes in shape are necessary for mitochondrial function and the production of new mitochondria. The OPA1 protein helps to regulate the shape of mitochondria by playing a key role in the fusion process.

The OPA1 protein is also involved in a process that takes place in mitochondria called oxidative phosphorylation, from which cells derive much of their energy. Additionally, the OPA1 protein plays a role in the maintenance of the DNA within mitochondria, called mitochondrial DNA (mtDNA), and in controlled cell death (apoptosis).

Health Conditions Related to Genetic Changes

Optic atrophy type 1

At least 240 mutations in the *OPA1* gene have been found to cause optic atrophy type 1. This condition typically results in vision loss beginning in childhood that worsens over time. Affected individuals may also have problems with color vision, particularly distinguishing between shades of blue and green.

Most of the *OPA1* gene mutations that cause optic atrophy type 1 create a premature stop signal in the instructions for making the OPA1 protein. As a result, an abnormally small protein is produced, which is likely to be unstable and broken down quickly. The most common mutation that causes optic atrophy type 1 in individuals of Danish ancestry results in an abnormally small protein by deleting one DNA building block (nucleotide) in the *OPA1* gene (written as 2826delT).

*OPA1* gene mutations that cause optic atrophy type 1 lead to problems in mitochondrial function. The mitochondria become misshapen and disorganized and have reduced energy-producing capabilities. The maintenance of mtDNA may also be impaired, resulting in mtDNA mutations that also contribute to mitochondrial dysfunction. Cells that contain these poorly functioning mitochondria are more susceptible to apoptosis. In particular, cells within the retina called retinal ganglion cells die over time. Specialized extensions of retinal ganglion cells, called axons, form the optic nerves, so when retinal ganglion cells die, the optic nerves break down...
(atrophy) and cannot transmit visual information to the brain. As the optic nerves atrophy, vision worsens, leading to the signs and symptoms of optic atrophy type 1. While the OPA1 protein is found in cells throughout the body, retinal ganglion cells appear to be particularly sensitive to the effects of OPA1 gene mutations. These cells have especially high energy requirements that make them more likely to malfunction and die when there are changes in mitochondrial function and decreases in energy production.

**Progressive external ophthalmoplegia**

**Other disorders**

About 20 percent of individuals with mutations in the OPA1 gene have the vision problems characteristic of optic atrophy type 1 (described above) with other health problems. Some OPA1 gene mutations cause a condition called optic atrophy type 1 and deafness, which results in both vision loss and hearing loss.

OPA1 mutations can also cause a condition known as autosomal dominant optic atrophy (ADOA)-plus syndrome. ADOA-plus syndrome involves vision and hearing loss, weakness in the muscles that control eye movement (progressive external ophthalmoplegia), difficulty with balance and coordination (ataxia), disturbances in the nerves used for muscle movement and sensation (motor and sensory neuropathy), and skeletal muscle weakness (myopathy).

The most severe condition cause by OPA1 gene mutations is Behr syndrome, which is characterized by neurological problems that begin by early childhood. Individuals with Behr syndrome develop optic atrophy, brain dysfunction (encephalopathy), loss of sensation and weakness in the limbs (peripheral neuropathy), difficulty coordinating movements (ataxia), feeding and digestive difficulties, and developmental delay.

The features of these conditions are likely caused by the loss of cells in multiple tissues due to poor mitochondrial function. It is unclear why OPA1 gene mutations affect only the eyes in individuals with optic atrophy type 1 but have more widespread effects in others. Researchers speculate that some OPA1 gene mutations lead to the production of an altered protein that interferes with the function of the normal protein produced from the non-mutated copy of the gene, further impairing OPA1 protein function.

While optic atrophy type 1 and ADOA-plus syndrome are caused by OPA1 gene mutations in one copy of the gene in each cell, individuals with Behr syndrome have mutations in both copies of the OPA1 gene in each cell. Alterations in both copies of the gene drastically reduce the amount of functional OPA1 protein, which likely leads to the severe signs and symptoms of Behr syndrome.
Chromosomal Location

Cytogenetic Location: 3q29, which is the long (q) arm of chromosome 3 at position 29

Molecular Location: base pairs 193,593,144 to 193,697,811 on chromosome 3 (Homo sapiens Annotation Release 109, GRCh38.p12) (NCBI)

Credit: Genome Decoration Page/NCBI

Other Names for This Gene

• dynamin-like 120 kDa protein, mitochondrial
• FLJ12460
• KIAA0567
• MGM1
• mitochondrial dynamin-like GTPase
• NPG
• NTG
• OPA1_HUMAN

Additional Information & Resources

Educational Resources

• eyeGENE: The National Ophthalmic Disease Genotyping and Phenotyping Network
  https://eyegene.nih.gov/
• Neuroscience (second edition, 2001): Central Projections of Retinal Ganglion Cells
  https://www.ncbi.nlm.nih.gov/books/NBK11145/
• Neuroscience (second edition, 2001): Central Projections of Retinal Ganglion Cells (image)
  https://www.ncbi.nlm.nih.gov/books/NBK11145/figure/A825/
• Washington University, St Louis: Neuromuscular Disease Center
  https://neuromuscular.wustl.edu/mitosyn.html#opa1

• Webvision--The Organization of the Retina and Visual System: Retinal Ganglion Cells

Clinical Information from GeneReviews
• Mitochondrial DNA Maintenance Defects Overview
  https://www.ncbi.nlm.nih.gov/books/NBK487393

• Optic Atrophy Type 1
  https://www.ncbi.nlm.nih.gov/books/NBK1248

Scientific Articles on PubMed
• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28OPA1%5BTIAB%5D%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last +720+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM
• OPA1 GENE
  http://omim.org/entry/605290

• OPTIC ATROPHY WITH OR WITHOUT DEAFNESS, OPHTHALMOPLEGIA, MYOPATHY, ATAXIA, AND NEUROPATHY
  http://omim.org/entry/125250

Research Resources
• Atlas of Genetics and Cytogenetics in Oncology and Haematology
  http://atlasgeneticsoncology.org/Genes/GC_OPA1.html

• ClinVar
  https://www.ncbi.nlm.nih.gov/clinvar?term=OPA1%5Bgene%5D

• HGNC Gene Symbol Report

• MITOchondrial DYNamics Variation Pages: OPA1 Gene Mutations
  http://www.mitodyn.org/home.php?select_db=OPA1

• Monarch Initiative
  https://monarchinitiative.org/gene/NCBIGene:4976
Sources for This Summary


  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/25796301

• OMIM: OPA1 GENE 
  http://omim.org/entry/605290

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17722006

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20157015 
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2842512/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18222991

Reprinted from Genetics Home Reference: 
https://ghr.nlm.nih.gov/gene/OPA1

Reviewed: August 2017
Published: May 14, 2019

Lister Hill National Center for Biomedical Communications
U.S. National Library of Medicine
National Institutes of Health
Department of Health & Human Services