

Autosomal dominant optic atrophy and cataract

Description

Autosomal dominant optic atrophy and cataract is an eye disorder that is characterized by impaired vision. Most affected individuals have decreased sharpness of vision (visual acuity) from birth, while others begin to experience vision problems in early childhood or later. In affected individuals, both eyes are usually affected equally. However, the severity of the vision loss varies widely, even among affected members of the same family, ranging from nearly normal vision to complete blindness.

Several abnormalities contribute to impaired vision in people with autosomal dominant optic atrophy and cataract. In the early stages of the condition, affected individuals experience a progressive loss of certain cells within the retina, which is a specialized light-sensitive tissue that lines the back of the eye. The loss of these cells (known as retinal ganglion cells) is followed by the degeneration (atrophy) of the nerves that relay visual information from the eyes to the brain (optic nerves), which contributes to vision loss. Atrophy of these nerves causes an abnormally pale appearance (pallor) of the optic nerves, which can be seen only during an eye examination. Most people with this disorder also have clouding of the lenses of the eyes (cataracts). This eye abnormality can develop anytime but typically appears in childhood. Other common eye problems in autosomal dominant optic atrophy and cataract include involuntary movements of the eyes (nystagmus), or problems with color vision (color vision deficiency) that make it difficult or impossible to distinguish between shades of blue and green.

Some people with autosomal dominant optic atrophy and cataract develop disturbances in the function of other nerves (neuropathy) besides the optic nerves. These disturbances can lead to problems with balance and coordination (cerebellar ataxia), an unsteady style of walking (gait), prickling or tingling sensations (paresthesias) in the arms and legs, progressive muscle stiffness (spasticity), or rhythmic shaking (tremors). In some cases, affected individuals have hearing loss caused by abnormalities of the inner ear (sensorineural deafness).

Frequency

Autosomal dominant optic atrophy and cataract is one form of autosomal dominant optic atrophy, a group of conditions that are estimated to affect 1 in 30,000 people worldwide, and approximately 1 in 10,000 people in Denmark. A form of optic atrophy called optic atrophy type 1 accounts for most cases, while autosomal dominant optic atrophy and

cataract is thought to represent only a few percent of autosomal dominant optic atrophy cases.

Causes

Autosomal dominant optic atrophy and cataract is caused by mutations in a gene called *OPA3*. The protein produced from this gene is made in cells and tissues throughout the body. The *OPA3* protein is found within mitochondria, which are the energy-producing centers of cells. While the exact function of the protein is unknown, it is thought to play a role in the organization of the shape and structure of the mitochondria and in controlled cell death (apoptosis).

Mutations in the *OPA3* gene lead to abnormal mitochondrial function. The mitochondria become misshapen and disorganized and have reduced energy-producing capabilities. Cells that contain these poorly functioning mitochondria seem to be more susceptible to apoptosis. In particular, affected cells that have high energy demands, such as retinal ganglion cells, are likely to die prematurely. Specialized extensions of retinal ganglion cells, called axons, form the optic nerves, so when retinal ganglion cells die, the optic nerves atrophy and cannot transmit visual information to the brain. Together, these effects reduce vision in affected individuals. It is likely that nerve cells in other parts of the body are similarly affected by dysfunctional mitochondria, resulting in the signs and symptoms of neuropathy in individuals with autosomal dominant optic atrophy and cataract. It is unclear how *OPA3* gene mutations lead to cataracts and other eye problems that can occur in autosomal dominant optic atrophy and cataracts.

[Learn more about the gene associated with Autosomal dominant optic atrophy and cataract](#)

- *OPA3*

Inheritance

This condition is inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to cause the disorder.

In most cases, an affected person has one parent with the condition.

Other Names for This Condition

- Autosomal dominant optic atrophy type 3
- *OPA3*
- *OPA3*, autosomal dominant
- Optic atrophy and cataract, autosomal dominant
- Optic atrophy type 3
- Optic atrophy, cataract, and neurologic disorder

Additional Information & Resources

Genetic Testing Information

- Genetic Testing Registry: Optic atrophy 3 (<https://www.ncbi.nlm.nih.gov/gtr/conditions/C1833809/>)

Genetic and Rare Diseases Information Center

- Autosomal dominant optic atrophy and cataract (<https://rarediseases.info.nih.gov/diseases/10203/index>)

Patient Support and Advocacy Resources

- National Organization for Rare Disorders (NORD) (<https://rarediseases.org/>)

Catalog of Genes and Diseases from OMIM

- OPTIC ATROPHY 3, AUTOSOMAL DOMINANT; OPA3 (<https://omim.org/entry/165300>)

Scientific Articles on PubMed

- PubMed (<https://pubmed.ncbi.nlm.nih.gov/?term=%28%28autosomal+dominant+optic+atrophy+and+cataract%5BTIAB%5D%29+OR+%28OPA3%5BTI%5D%29%29+AND+english%5BIa%5D+AND+human%5Bmh%5D>)

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Last updated February 1, 2019