CACNA1F gene
calcium voltage-gated channel subunit alpha1 F

Normal Function
The \textit{CACNA1F} gene belongs to a family of genes that provide instructions for making calcium channels. These channels, which transport positively charged calcium atoms (calcium ions) across cell membranes, play a key role in a cell's ability to generate and transmit electrical signals.

The \textit{CACNA1F} gene provides instructions for making one part (the alpha-1 subunit) of a calcium channel called CaV1.4. This subunit forms the hole (pore) in the cell membrane through which calcium ions can flow. CaV1.4 channels are found in many types of cells, although they play a particularly important role in a specialized tissue at the back of the eye called the retina. Within the retina, the channels are located in light-detecting cells called photoreceptors. The retina contains two types of photoreceptors: rods and cones. Rods are responsible for vision in low light. Cones provide vision in bright light, including color vision.

CaV1.4 channels appear to play a critical role in normal vision. Studies suggest they help relay visual signals from rods and cones to other retinal cells called bipolar cells. This signaling is an essential step in the transmission of visual information from the eyes to the brain.

Health Conditions Related to Genetic Changes

\textbf{X-linked congenital stationary night blindness}

More than 70 mutations in the \textit{CACNA1F} gene have been identified in people with X-linked congenital stationary night blindness. Mutations in this gene are responsible for the incomplete form of the disorder, which is characterized by vision problems including a loss of sharpness (reduced acuity), nearsightedness (myopia), involuntary movements of the eyes (nystagmus), and eyes that do not look in the same direction (strabismus). Many affected individuals also have difficulty seeing in low light (night blindness).

\textit{CACNA1F} mutations change the structure of the alpha-1 subunit, which alters or eliminates the function of CaV1.4 channels. These changes prevent the normal transport of calcium ions across the cell membrane of photoreceptor cells. A loss of calcium ion transport disrupts the ability of both rods and cones to transmit visual signals, which impairs vision.

\textbf{Cone-rod dystrophy}
Other disorders

Mutations in the CACNA1F gene cause several other rare disorders that impair vision. These include Åland Island eye disease, X-linked cone-rod dystrophy, an X-linked retinal disorder in New Zealand, and retinal and optic disc atrophy. Each of these disorders has been reported in only a few individuals or families worldwide. They cause vision problems similar to those of X-linked congenital stationary night blindness.

Researchers have identified at least one CACNA1F mutation that can cause Åland Island eye disease (also known as Forsius-Eriksson syndrome). This condition was first described in a family from the Åland Islands, which are in the Baltic Sea off the coast of Sweden. Åland Island eye disease is characterized by reduced visual acuity, nystagmus, an irregular curvature of the front part of the eye (astigmatism), myopia, abnormal color vision, and night blindness. The mutation associated with this disorder deletes a segment of genetic material from the CACNA1F gene, which significantly alters the structure of the alpha-1 subunit of CaV1.4 channels. These changes prevent the normal transport of calcium ions across the cell membrane of photoreceptor cells. A loss of calcium ion transport disrupts the ability of both rods and cones to transmit visual signals.

At least one other CACNA1F mutation is responsible for X-linked cone-rod dystrophy (also known as CORDX3). The signs and symptoms of this condition include reduced visual acuity, an increased sensitivity to light (photophobia), myopia, and impaired color vision. These vision problems tend to worsen over time. The mutation associated with this disorder deletes part of the alpha-1 subunit, which likely prevents the production of functional CaV1.4 channels. A loss of these channels keeps photoreceptor cells from relaying visual signals normally, which leads to impaired vision.

A CACNA1F mutation has also been found to cause an X-linked retinal disorder in a large Maori family from New Zealand. The major features of this disorder include reduced visual acuity, abnormal color vision, photophobia, and mild nystagmus. Some affected individuals have also had intellectual disability. The CACNA1F mutation associated with this condition alters a single protein building block (amino acid) in the alpha-1 subunit, which appears to overactivate CaV1.4 channels. The resulting increase in calcium ion transport probably disrupts the transmission of visual signals in the retina.

Another eye disorder, known as retinal and optic disc atrophy, has been associated with a CACNA1F mutation in two Japanese brothers. The affected individuals experienced a progressive decline in visual acuity and color vision. These vision problems were caused by deterioration of the retina, including an area called the optic disc (which is where the retina connects with the nerve that relays visual information to the brain). The CACNA1F mutation responsible for retinal and optic disc atrophy alters the structure of the alpha-1 subunit, which probably leads to the production of nonfunctional CaV1.4 channels. This mutation has also been identified
in at least one Japanese family with X-linked congenital stationary night blindness. It is unclear why this single genetic change can cause different vision abnormalities in different families.

**Chromosomal Location**

Cytogenetic Location: Xp11.23, which is the short (p) arm of the X chromosome at position 11.23

Molecular Location: base pairs 49,205,063 to 49,233,404 on the X chromosome (Homo sapiens Updated Annotation Release 109.20200522, GRCh38.p13) (NCBI)

![Chromosomal Location Diagram](image)

Credit: Genome Decoration Page/NCBI

**Other Names for This Gene**

- AIED
- CAC1F_HUMAN
- calcium channel, voltage-dependent, L type, alpha 1F subunit
- Cav1.4
- Cav1.4alpha1
- COD3
- COD4
- CORDX
- CORDX3
- CSNB2
- CSNB2A
- CSNBX2
- JM8
- JMC8
- OA2
Additional Information & Resources

Educational Resources

• Madame Curie Bioscience Database: α1F/ CaV1.4  
  https://www.ncbi.nlm.nih.gov/books/NBK6181/#A30881

• Neuromuscular Disease Center, Washington University: Calcium channels  
  https://neuromuscular.wustl.edu/mother/chan.html#ca

Clinical Information from GeneReviews

• X-Linked Congenital Stationary Night Blindness  
  https://www.ncbi.nlm.nih.gov/books/NBK1245

Scientific Articles on PubMed

• PubMed  
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28CACNA1F%5BTIAB%5D%29+OR+%28Cav1.4%5BTIAB%5D%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+2160+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

• ALAND ISLAND EYE DISEASE  
  http://omim.org/entry/300600

• CALCIUM CHANNEL, VOLTAGE-DEPENDENT, ALPHA-1F SUBUNIT  
  http://omim.org/entry/300110

• CONE-ROD DYSTROPHY, X-LINKED, 3  
  http://omim.org/entry/300476

Research Resources

• Atlas of Genetics and Cytogenetics in Oncology and Haematology  
  http://atlasgeneticsoncology.org/Genes/GC_CACNA1F.html

• ClinVar  
  https://www.ncbi.nlm.nih.gov/clinvar?term=CACNA1F%5Bgene%5D

• HGNC Gene Symbol Report  

• Monarch Initiative  
  https://monarchinitiative.org/gene/NCBIGene:778

• NCBI Gene  

• UniProt  
  https://www.uniprot.org/uniprot/O60840
Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/9662400

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/11281458

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19151588

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15634789

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16476079

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15807819

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17525176

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16505158 
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2564595/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/14973233
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12860808

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17949918

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/9662399

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12111638

Reprinted from Genetics Home Reference:

Reviewed: May 2009
Published: June 23, 2020

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