PSEN2 gene
presenilin 2

Normal Function

The *PSEN2* gene provides instructions for making a protein called presenilin 2. Presenilin 2 helps process proteins that transmit chemical signals from the cell membrane into the nucleus. Once in the nucleus, these signals turn on (activate) genes that are important for cell growth and maturation.

Presenilin 2 is best known for its role in processing amyloid precursor protein, which is found in the brain and other tissues. Research suggests that presenilin 2 works together with other enzymes to cut amyloid precursor protein into smaller segments (peptides). One of these peptides is called soluble amyloid precursor protein (sAPP), and another is called amyloid beta peptide. Recent evidence suggests that sAPP has growth-promoting properties and may play a role in the formation of neurons in the brain both before and after birth. Other functions of sAPP and amyloid beta peptide are under investigation.

Health Conditions Related to Genetic Changes

**Alzheimer disease**

At least 11 mutations in the *PSEN2* gene have been shown to cause early-onset Alzheimer disease. Mutations in this gene account for less than 5 percent of all early-onset cases of the disorder.

Two of the most common *PSEN2* mutations that cause early-onset Alzheimer disease change single protein building blocks (amino acids) used to make presenilin 2. One mutation replaces the amino acid asparagine with the amino acid isoleucine at position 141 (written as Asn141Ile or N141I). The other mutation changes the amino acid methionine to the amino acid valine at position 239 (written as Met239Val or M239V). These mutations appear to disrupt the processing of amyloid precursor protein, leading to the overproduction of amyloid beta peptide. This protein fragment can build up in the brain and form clumps called amyloid plaques that are characteristic of Alzheimer disease. A buildup of toxic amyloid beta peptide and amyloid plaques may lead to the death of neurons and the progressive signs and symptoms of this disorder.

**Familial dilated cardiomyopathy**
Chromosomal Location

Cytogenetic Location: 1q42.13, which is the long (q) arm of chromosome 1 at position 42.13

Molecular Location: base pairs 226,870,572 to 226,903,829 on chromosome 1 (Homo sapiens Updated Annotation Release 109.20190607, GRCh38.p13) (NCBI)

Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- AD3-like protein
- AD3L
- AD3LP
- AD4
- AD5
- Alzheimer's disease 3-like
- E5-1
- presenilin 2 (Alzheimer disease 4)
- PS2 protein (alzheimer-associated)
- PSN2_HUMAN
- PSNL2
- STM2

Additional Information & Resources

Educational Resources

Clinical Information from GeneReviews

- Alzheimer Disease Overview

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28PSEN2%5BTIAB%5D%29+OR+%28presenilin+2%5BTIAB%5D%29%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%28Blia%5D+AND+human%5Bmh%5D+AND+%22last+1080+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- PRESENILIN 2
  http://omim.org/entry/600759

Research Resources

- Alzheimer Disease & Frontotemporal Dementia Mutation Database

- Alzheimer Research Forum: AlzGene database

- Atlas of Genetics and Cytogenetics in Oncology and Haematology
  http://atlasgeneticsoncology.org/Genes/PSEN2ID41883ch1q42.html

- ClinVar

- HGNC Gene Symbol Report

- Monarch Initiative
  https://monarchinitiative.org/gene/NCBIGene:5664

- NCBI Gene

- UniProt
  https://www.uniprot.org/uniprot/P49810
Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15304236

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