MPZ gene
myelin protein zero

Normal Function

The MPZ gene provides instructions for making a protein called myelin protein zero. It is the most abundant protein in myelin, a protective substance that covers nerves and promotes the efficient transmission of nerve impulses. Myelin protein zero is produced by specialized cells called Schwann cells, which wrap around and insulate peripheral nerves. Peripheral nerves connect the brain and spinal cord to muscles and to sensory cells that detect sensations such as touch, pain, heat, and sound. Myelin protein zero is required for the proper formation and maintenance of myelin. This protein is an adhesion molecule, which means it acts like molecular glue. It plays a role in tightly packing the myelin around nerve cells (myelin compaction).

Health Conditions Related to Genetic Changes

Charcot-Marie-Tooth disease

Researchers have identified more than 120 MPZ gene mutations that cause a form of Charcot-Marie-Tooth disease known as type 1B. Charcot-Marie-Tooth syndrome is a disorder characterized by muscle weakness and sensory problems, especially in the hands and feet. Many of the MPZ gene mutations that cause Charcot-Marie-Tooth syndrome change single protein building blocks (amino acids) in myelin protein zero. Other mutations lead to a protein that is missing one or more amino acids. The altered myelin protein zero probably cannot interact properly with other myelin components, which may disrupt myelin compaction. As a result, peripheral nerves cannot trigger muscle movement or relay information from sensory cells back to the brain, leading to the weakness and sensory problems characteristic of Charcot-Marie-Tooth disease.

Some MPZ gene mutations cause a severe form of type 1B Charcot-Marie-Tooth disease. Symptoms begin during infancy or early childhood and include delayed development of motor skills such as walking. This form of Charcot-Marie-Tooth disease is sometimes called Dejerine-Sottas syndrome, congenital hypomyelination, or Roussy-Levy syndrome. Researchers believe that the MPZ gene mutations that cause the severe form of the disorder probably disrupt the formation of myelin during early development.

Several mutations in the MPZ gene cause other forms of Charcot-Marie-Tooth disease known as type 2I, type 2J, and dominant intermediate D. These forms of Charcot-Marie-Tooth disease, which often do not become evident until adulthood, affect the specialized outgrowths from nerve cells (axons) that transmit impulses to
muscles and other nerve cells. People with type 2J Charcot-Marie-Tooth disease may also have hearing loss and abnormalities in the opening of the eye through which light passes (the pupil). It is unclear how MPZ gene mutations cause these abnormalities.

Chromosomal Location

Cytogenetic Location: 1q23.3, which is the long (q) arm of chromosome 1 at position 23.3

Molecular Location: base pairs 161,303,594 to 161,309,972 on chromosome 1 (Homo sapiens Annotation Release 109, GRCh38.p12) (NCBI)

Credit: Genome Decoration Page/NCBI

Other Names for This Gene

• CMT1B
• CMT2I
• CMT2J
• HMSN1B
• MPP
• myelin glycoprotein P-zero
• myelin peripheral protein
• myelin protein zero (Charcot-Marie-Tooth neuropathy 1B)
• MYP0_HUMAN
• P0 Glycoprotein
• P0 Protein
Additional Information & Resources

Educational Resources

• Basic Neurochemistry (sixth edition, 1999): Cell Adhesion Molecules in Myelination
  https://www.ncbi.nlm.nih.gov/books/NBK28158/

• Basic Neurochemistry (sixth edition, 1999): The Myelin Sheath
  https://www.ncbi.nlm.nih.gov/books/NBK27954/

Clinical Information from GeneReviews

• Charcot-Marie-Tooth Hereditary Neuropathy Overview
  https://www.ncbi.nlm.nih.gov/books/NBK1358

Scientific Articles on PubMed

• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28MPZ%5BTIAB%5D%29%29+OR+%28myelin+protein+zero%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+1440+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

• MYELIN PROTEIN ZERO
  http://omim.org/entry/159440

Research Resources

• ClinVar

• HGNC Gene Family: Ig-like cell adhesion molecule family
  https://www.genenames.org/cgi-bin/genefamilies/set/1410

• HGNC Gene Family: V-set domain containing
  https://www.genenames.org/cgi-bin/genefamilies/set/590

• HGNC Gene Symbol Report

• Inherited Peripheral Neuropathies Mutation Database
  http://www.molgen.ua.ac.be/CMTMutations/Mutations/Mutations.cfm?Context=2

• Monarch Initiative
Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16856148

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20301532

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/11835375

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/25625223

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18337304

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15298082

- OMIM: MYELIN PROTEIN ZERO 
  http://omim.org/entry/159440

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19293842 
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2986589/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16775378

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12402337
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16414078

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/8816708

Reprinted from Genetics Home Reference:  

Reviewed: October 2018  
Published: October 23, 2018

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