Hereditary neuropathy with liability to pressure palsies

Hereditary neuropathy with liability to pressure palsies is a disorder that affects peripheral nerves. These nerves connect the brain and spinal cord to muscles and sensory cells that detect touch, pain, and temperature. In people with this disorder, the peripheral nerves are unusually sensitive to pressure, such as the pressure that occurs when carrying heavy grocery bags, leaning on an elbow, or sitting without changing position, particularly with crossed legs. These activities would not normally cause sensation problems in people without the disorder.

Hereditary neuropathy with liability to pressure palsies is characterized by recurrent episodes of numbness, tingling, and loss of muscle function (palsy) in the region associated with the affected nerve, usually an arm, hand, leg, or foot. An episode can last from several minutes to several months, but recovery is usually complete. Repeated incidents, however, can cause permanent muscle weakness or loss of sensation. This disorder is also associated with pain in the limbs, especially the hands.

A pressure palsy episode results from pressure on a single nerve, and any peripheral nerve can be affected. Although episodes often recur, they can affect different nerves. The most common problem sites involve nerves in the wrists, elbows, and knees. The fingers, shoulders, hands, feet, and scalp can also be affected. Many people with this disorder experience carpal tunnel syndrome, which occurs when a nerve in the wrist (the median nerve) is involved. Carpal tunnel syndrome is characterized by numbness, tingling, and weakness in the hand and fingers. An episode in the hand may affect fine motor activities such as writing, opening jars, and fastening buttons. An episode of nerve compression in the knee can lead to a condition called foot drop, which makes walking, climbing stairs, or driving difficult or impossible.

The symptoms of hereditary neuropathy with liability to pressure palsies usually begin during adolescence or early adulthood but may develop anytime from childhood to late adulthood. Symptoms vary in severity; many people never realize they have the disorder, while some people experience prolonged disability. Hereditary neuropathy with liability to pressure palsies does not affect life expectancy.

Frequency

Hereditary neuropathy with liability to pressure palsies is estimated to occur in 2 to 5 per 100,000 individuals.

Causes

Hereditary neuropathy with liability to pressure palsies is most commonly caused by the deletion of one copy of the PMP22 gene. This loss reduces the amount of PMP22 gene product, which normally serves an important role in maintaining the health of peripheral nerves. Other genetic and environmental factors can also contribute to the development of this condition.
protein produced. Other PMP22 gene mutations that lead to a reduction of PMP22 protein can also cause the condition.

The PMP22 protein is a component of myelin, a protective substance that covers nerves and promotes the efficient transmission of nerve impulses. Studies suggest that the PMP22 protein is particularly important in protecting nerves from physical pressure, helping them restore their structure after compression. Compression can interrupt nerve signaling, leading to the sensation commonly referred to as a limb "falling asleep." The ability of nerves to recover from normal, day-to-day pressure, for example when sitting for long periods, keeps the limbs from constantly losing sensation.

The consequences of PMP22 gene mutations are not clearly understood. A shortage of PMP22 protein appears to make nerves less able to recover from compression, which impairs the transmission of nerve impulses, causing the signs and symptoms of hereditary neuropathy with liability to pressure palsies.

Inheritance Pattern

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.

Other Names for This Condition

- compression neuropathy
- entrapment neuropathy
- familial pressure sensitive neuropathy
- hereditary motor and sensory neuropathy
- hereditary pressure sensitive neuropathy
- HNPP
- inherited tendency to pressure palsies
- tomaculous neuropathy

Diagnosis & Management

Genetic Testing

- Genetic Testing Registry: Hereditary liability to pressure palsies

Other Diagnosis and Management Resources

- GeneReview: Hereditary Neuropathy with Liability to Pressure Palsies
  https://www.ncbi.nlm.nih.gov/books/NBK1392
- MedlinePlus Encyclopedia: Carpal Tunnel Syndrome
  https://medlineplus.gov/ency/article/000433.htm
General Information from MedlinePlus

- Diagnostic Tests
  https://medlineplus.gov/diagnostictests.html
- Drug Therapy
  https://medlineplus.gov/drugtherapy.html
- Genetic Counseling
  https://medlineplus.gov/geneticcounseling.html
- Palliative Care
  https://medlineplus.gov/palliativecare.html
- Surgery and Rehabilitation
  https://medlineplus.gov/surgeryandrehabilitation.html

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Carpal Tunnel Syndrome
  https://medlineplus.gov/ency/article/000433.htm
- Encyclopedia: Peripheral Neuropathy
  https://medlineplus.gov/ency/article/000593.htm
- Health Topic: Carpal Tunnel Syndrome
  https://medlineplus.gov/carpaltunnelsyndrome.html
- Health Topic: Peripheral Nerve Disorders
  https://medlineplus.gov/peripheralnervedisorders.html

Genetic and Rare Diseases Information Center

- Hereditary neuropathy with liability to pressure palsy

Educational Resources

- Johns Hopkins Medicine: Peripheral Nerve System
  https://www.hopkinsmedicine.org/neurology_neurosurgery/centers_clinics/peripheral_nerve_surgery/conditions/peripheral_nerve_system.html
- MalaCards: hereditary neuropathy with liability to pressure palsy
  http://www.malacards.org/card/hereditary_neuropathy_with_liability_to_pressure_palsy
- Merck Manual Consumer Version
• National Health Service (UK)
  https://www.nhs.uk/conditions/hereditary-neuropathy/

• Orphanet: Hereditary neuropathy with liability to pressure palsies
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=640

• University of Chicago Center for Peripheral Neuropathy
  http://peripheralneuropathycenter.uchicago.edu/learnaboutpn/typesofpn/hereditary/hnpp.shtml

Patient Support and Advocacy Resources

• Charcot-Marie-Tooth Association
  http://www.cmtausa.org/understanding-cmt/types-and-causes/more-about-hnpp/

• The Foundation for Peripheral Neuropathy
  https://www.foundationforpn.org/

Clinical Information from GeneReviews

• Hereditary Neuropathy with Liability to Pressure Palsies
  https://www.ncbi.nlm.nih.gov/books/NBK1392

Research Studies from ClinicalTrials.gov

• ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22hereditary+neuropathy+with+liability+to+pressure+palsies%22

Scientific Articles on PubMed

• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28Hereditary+Motor+and+Sensory+Neuropathies%5BMAJR%5D%29+AND+%28%28hnpp%5BTIAB%5D%29+OR+%28tomaculous+neuropathy%5BTIAB%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1800+days%22+AND+last+1800+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

• NEUROPATHY, HEREDITARY, WITH LIABILITY TO PRESSURE PALSIES
  http://omim.org/entry/162500
Sources for This Summary

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  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3676309/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20301566

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16106622

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16775374

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