Limb-girdle muscular dystrophy

Limb-girdle muscular dystrophy is a term for a group of diseases that cause weakness and wasting of the muscles in the arms and legs. The muscles most affected are those closest to the body (proximal muscles), specifically the muscles of the shoulders, upper arms, pelvic area, and thighs.

The severity, age of onset, and features of limb-girdle muscle dystrophy vary among the many subtypes of this condition and may be inconsistent even within the same family. Signs and symptoms may first appear at any age and generally worsen with time, although in some cases they remain mild.

In the early stages of limb-girdle muscular dystrophy, affected individuals may have an unusual walking gait, such as waddling or walking on the balls of their feet, and may also have difficulty running. They may need to use their arms to press themselves up from a squatting position because of their weak thigh muscles. As the condition progresses, people with limb-girdle muscular dystrophy may eventually require wheelchair assistance.

Muscle wasting may cause changes in posture or in the appearance of the shoulder, back, and arm. In particular, weak shoulder muscles tend to make the shoulder blades (scapulae) "stick out" from the back, a sign known as scapular winging. Affected individuals may also have an abnormally curved lower back (lordosis) or a spine that curves to the side (scoliosis). Some develop joint stiffness (contractures) that can restrict movement in their hips, knees, ankles, or elbows. Overgrowth (hypertrophy) of the calf muscles occurs in some people with limb-girdle muscular dystrophy.

Weakening of the heart muscle (cardiomyopathy) occurs in some forms of limb-girdle muscular dystrophy. Some affected individuals experience mild to severe breathing problems related to the weakness of muscles needed for breathing. In some cases, the breathing problems are severe enough that affected individuals need to use a machine to help them breathe (mechanical ventilation).

Intelligence is generally unaffected in limb-girdle muscular dystrophy; however, developmental delay and intellectual disability have been reported in rare forms of the disorder.

Frequency

It is difficult to determine the prevalence of limb-girdle muscular dystrophy because its features vary and overlap with those of other muscle disorders. Prevalence estimates range from 1 in 14,500 to 1 in 123,000 individuals.
Causes

The various forms of limb-girdle muscular dystrophy are caused by mutations in many different genes. These genes provide instructions for making proteins that are involved in muscle maintenance and repair.

Some of the proteins produced from these genes assemble with other proteins into larger protein complexes. These complexes maintain the physical integrity of muscle tissue and allow the muscles to contract. Other proteins participate in cell signaling, cell membrane repair, or the removal of potentially toxic wastes from muscle cells.

Limb-girdle muscular dystrophy is classified based on its inheritance pattern and genetic cause. Limb-girdle muscular dystrophy type 1 includes forms of the disorder that have an inheritance pattern called autosomal dominant. Mutations in the LMNA gene cause limb-girdle muscular dystrophy type 1B. Limb-girdle muscular dystrophy type 1C is one of a group of muscle disorders called caveolinopathies caused by mutations in the CAV3 gene.

Limb-girdle muscular dystrophy type 2 includes forms of the disorder that have an inheritance pattern called autosomal recessive. Calpainopathy, or limb-girdle muscular dystrophy type 2A, is caused by mutations in the CAPN3 gene. Type 2A is the most common form of limb-girdle muscular dystrophy, accounting for about 30 percent of cases. Dysferlinopathy, also called limb-girdle muscular dystrophy type 2B, is caused by mutations in the DYSF gene.

Sarcoglycanopathies are forms of limb-girdle muscular dystrophy caused by mutations in the SGCA, SGCB, SGCG, and SGCD genes. These sarcoglycanopathies are known as limb-girdle muscular dystrophy types 2D, 2E, 2C, and 2F respectively.

A TTN gene mutation causes limb-girdle muscular dystrophy type 2J, which has been identified only in the Finnish population. Mutations in the ANOS gene cause limb-girdle muscular dystrophy type 2L. Mutations in several other genes cause forms of limb-girdle muscular dystrophy called dystroglycanopathies, including limb-girdle muscular dystrophy types 2I, 2K, 2M, and 2N.

Other rare forms of limb-girdle muscular dystrophy are caused by mutations in several other genes, some of which have not been identified.

Inheritance Pattern

Limb-girdle muscular dystrophy can have different inheritance patterns.

Most forms of this condition are inherited in an autosomal recessive pattern, which means both copies of the gene in each cell have mutations. The parents of an individual with an autosomal recessive condition each carry one copy of the mutated gene, but they typically do not show signs and symptoms of the condition.

Several rare forms of limb-girdle muscular dystrophy are 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

- LGMD
- limb-girdle syndrome
- myopathic limb-girdle syndrome

Diagnosis & Management

Genetic Testing Information

- What is genetic testing?/primer/testing/genetictesting
- Genetic Testing Registry: Limb-girdle muscular dystrophy

Research Studies from ClinicalTrials.gov

- ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22limb-girdle+muscular+dystrophy%22

Other Diagnosis and Management Resources

- Johns Hopkins Medicine
  https://www.hopkinsmedicine.org/neurology_neurosurgery/centers_clinics/muscular_dystrophy/conditions/limb_girdle_muscular_dystrophy.html

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Limb-Girdle Muscular Dystrophies
  https://medlineplus.gov/ency/article/000711.htm
- Health Topic: Muscular Dystrophy
  https://medlineplus.gov/musculardystrophy.html

Genetic and Rare Diseases Information Center

- Limb-girdle muscular dystrophy

Additional NIH Resources

- National Institute of Neurological Disorders and Stroke (NINDS): Muscular Dystrophy
  https://www.ninds.nih.gov/Disorders/All-Disorders/Muscular-Dystrophy-Information-Page
Educational Resources

• MalaCards: limb-girdle muscular dystrophy
  https://www.malacards.org/card/limb_girdle_muscular_dystrophy

• Orphanet: Limb-girdle muscular dystrophy
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=263

• University of Washington Neuromuscular Disease Center
  https://neuromuscular.wustl.edu/musdist/lg.html

Patient Support and Advocacy Resources

• Jain Foundation
  http://www.jain-foundation.org/

• Muscular Dystrophy Association

• Muscular Dystrophy Association New Zealand

• Muscular Dystrophy Australia

• Muscular Dystrophy Canada

• Muscular Dystrophy UK
  https://www.musculardystrophyuk.org/about-muscle-wasting-conditions/limb-girdle-muscular-dystrophy/

• National Organization for Rare Disorders (NORD)
  https://rarediseases.org/rare-diseases/limb-girdle-muscular-dystrophies/

Clinical Information from GeneReviews

• Calpainopathy
  https://www.ncbi.nlm.nih.gov/books/NBK1313

• Caveolinopathies
  https://www.ncbi.nlm.nih.gov/books/NBK1385

• Dysferlinopathy
  https://www.ncbi.nlm.nih.gov/books/NBK1303
Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28Muscular+Dystrophies,+Limb-Girdle%5BMAJR%5D%29+AND+%28limb-girdle+muscular+dystrophy%5BTIAB %5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last +720+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- MUSCULAR DYSTROPHY-DYSTROGLYCANOPATHY (LIMB-GIRDLE), TYPE C, 1
  http://omim.org/entry/609308
- MUSCULAR DYSTROPHY-DYSTROGLYCANOPATHY (LIMB-GIRDLE), TYPE C, 2
  http://omim.org/entry/613158
- MUSCULAR DYSTROPHY-DYSTROGLYCANOPATHY (LIMB-GIRDLE), TYPE C, 3
  http://omim.org/entry/613157
- MUSCULAR DYSTROPHY-DYSTROGLYCANOPATHY (LIMB-GIRDLE), TYPE C, 4
  http://omim.org/entry/611588
- MUSCULAR DYSTROPHY-DYSTROGLYCANOPATHY (LIMB-GIRDLE), TYPE C, 5
  http://omim.org/entry/607155
- MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL DOMINANT 1
  http://omim.org/entry/603511
- MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL DOMINANT 2
  http://omim.org/entry/608423
- MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL DOMINANT 3
  http://omim.org/entry/609115
- MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL RECESSIVE 1
  http://omim.org/entry/253600
- MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL RECESSIVE 2
  http://omim.org/entry/253601
- MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL RECESSIVE 3
  http://omim.org/entry/608099
- MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL RECESSIVE 4
  http://omim.org/entry/604286
- MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL RECESSIVE 5
  http://omim.org/entry/253700
• MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL RECESSIVE 6
  http://omim.org/entry/601287
• MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL RECESSIVE 7
  http://omim.org/entry/601954
• MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL RECESSIVE 8
  http://omim.org/entry/254110
• MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL RECESSIVE 10
  http://omim.org/entry/608807
• MUSCULAR DYSTROPHY, LIMB-GIRDLE, AUTOSOMAL RECESSIVE 12
  http://omim.org/entry/611307
• MUSCULAR DYSTROPHY, LIMB-GIRDLE, TYPE 1A
  http://omim.org/entry/159000
• MUSCULAR DYSTROPHY, LIMB-GIRDLE, TYPE 1B
  http://omim.org/entry/159001
• MUSCULAR DYSTROPHY, LIMB-GIRDLE, TYPE 1C
  http://omim.org/entry/607801
• MUSCULAR DYSTROPHY, LIMB-GIRDLE, TYPE 1H
  http://omim.org/entry/613530

Medical Genetics Database from MedGen
• Limb-girdle muscular dystrophy

Sources for This Summary
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/21150381
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19584897
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2987183/
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  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16934440

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15043707

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19019316
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4514705/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19781108
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3279956/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17027860

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18974555

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