Diastrophic dysplasia

Diastrophic dysplasia is a disorder of cartilage and bone development. Affected individuals have short stature with very short arms and legs. Most also have early-onset joint pain (osteoarthritis) and joint deformities called contractures, which restrict movement. These joint problems often make it difficult to walk and tend to worsen with age. Additional features of diastrophic dysplasia include an inward- and upward-turning foot (clubfoot), progressive abnormal curvature of the spine, and unusually positioned thumbs (hitchhiker thumbs). About half of infants with diastrophic dysplasia are born with an opening in the roof of the mouth (a cleft palate). Swelling of the external ears is also common in newborns and can lead to thickened, deformed ears.

The signs and symptoms of diastrophic dysplasia are similar to those of another skeletal disorder called atelosteogenesis type 2; however, diastrophic dysplasia tends to be less severe. Although some affected infants have breathing problems, most people with diastrophic dysplasia live into adulthood.

Frequency

Although the exact prevalence of diastrophic dysplasia is unknown, researchers estimate that it affects about 1 in 500,000 newborns in the United States. This condition is more common in Finland, where it affects about 1 in 33,000 newborns.

Causes

Diastrophic dysplasia is one of several skeletal disorders caused by mutations in the SLC26A2 gene. This gene provides instructions for making a protein that is essential for the normal development of cartilage and for its conversion to bone. Cartilage is a tough, flexible tissue that makes up much of the skeleton during early development. Most cartilage is later converted to bone, except for the cartilage that continues to cover and protect the ends of bones and is present in the nose and external ears. Mutations in the SLC26A2 gene alter the structure of developing cartilage, preventing bones from forming properly and resulting in the skeletal problems characteristic of diastrophic dysplasia.

Inheritance Pattern

This condition is 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.
Other Names for This Condition
- Diastrophic dwarfism
- DTD

Diagnosis & Management
Genetic Testing Information
- What is genetic testing?
  /primer/testing/genetictesting
- Genetic Testing Registry: Diastrophic dysplasia

Other Diagnosis and Management Resources
- GeneReview: Diastrophic Dysplasia
  https://www.ncbi.nlm.nih.gov/books/NBK1350

Additional Information & Resources
Health Information from MedlinePlus
- Health Topic: Bone Diseases
  https://medlineplus.gov/bonediseases.html
- Health Topic: Dwarfism
  https://medlineplus.gov/dwarfism.html

Genetic and Rare Diseases Information Center
- Diastrophic dysplasia
  https://rarediseases.info.nih.gov/diseases/6275/diastrophic-dysplasia

Additional NIH Resources
- National Institute of Arthritis and Musculoskeletal and Skin Diseases: Heritable Disorders of Connective Tissue
  https://www.niams.nih.gov/health-topics/heritable-disorders-connective-tissue

Educational Resources
- Children's Hospital of Philadelphia
  https://www.chop.edu/conditions-diseases/diastrophic-dysplasia
- Kaiser Permanente Genetics Northern California
  https://mydoctor.kaiserpermanente.org/ncal/specialty/genetics/resources/conditions/diastrophic_dystrophy.jsp
- KidsHealth from the Nemours Foundation
• MalaCards: diastrophic dysplasia
  https://www.malacards.org/card/diastrophic_dysplasia

• Nemours Children’s Health System
  https://www.nemours.org/services/skeletal-dysplasia/diastrophic.html?tab=about

• Orphanet: Diastrophic dwarfism
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=628

Patient Support and Advocacy Resources

• European Skeletal Dysplasia Network
  http://www.esdn.org/

• Human Growth Foundation
  http://hgfound.org/

• Little People of America
  https://www.lpaonline.org/

• Little People UK
  https://littlepeopleuk.org/

• National Organization for Rare Disorders
  https://rarediseases.org/rare-diseases/diastrophic-dysplasia/

• Resource list from the University of Kansas Medical Center
  http://www.kumc.edu/gec/support/dwarfism.html

• The MAGIC Foundation
  https://www.magicfoundation.org/

Clinical Information from GeneReviews

• Diastrophic Dysplasia
  https://www.ncbi.nlm.nih.gov/books/NBK1350

Scientific Articles on PubMed

• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28diastrophic+dysplasia%5BTIAB%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1440+days%22+AND+human%5Bdp%5D

Catalog of Genes and Diseases from OMIM

• DIASTROPHIC DYSPLASIA
  http://omim.org/entry/222600
Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20301524

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17602446

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/23657516

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15308906

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/11241838

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