Camurati-Engelmann disease

Camurati-Engelmann disease is a skeletal condition that is characterized by abnormally thick bones (hyperostosis) in the arms, legs, and skull.

The thick limb bones can lead to bone pain and muscle weakness in the arms and legs and cause individuals with Camurati-Engelmann disease to tire quickly. Bone pain ranges from mild to severe and can increase with stress, activity, or cold weather. Leg weakness can make it difficult to stand up from a seated position and some affected individuals develop a waddling or unsteady walk. Additional limb abnormalities include joint deformities (contractures), knock knees (a condition in which the lower legs are positioned at an outward angle), and flat feet (pes planus). Swelling and redness (erythema) of the limbs and an abnormal curvature of the spine can also occur.

Individuals with Camurati-Engelmann disease may have an unusually thick skull, which can lead to an abnormally large head (macrocephaly) and lower jaw (mandible), a prominent forehead (frontal bossing), and bulging eyes with shallow eye sockets (ocular proptosis). These changes to the head and face become more prominent with age and are most noticeable in affected adults. In about a quarter of individuals with Camurati-Engelmann disease, the thickened skull increases pressure on the brain or compresses the spinal cord, which can cause a variety of neurological problems, including headaches, hearing loss, vision problems, dizziness (vertigo), ringing in the ears (tinnitus), and facial paralysis.

The degree of hyperostosis varies among individuals with Camurati-Engelmann disease as does the age at which they experience their first symptoms.

Other, rare features of Camurati-Engelmann disease include abnormally long limbs in proportion to height, a decrease in muscle mass and body fat, delayed teething (dentition), frequent cavities, delayed puberty, a shortage of red blood cells (anemia), an enlarged liver and spleen (hepatosplenomegaly), thinning of the skin, and excessively sweaty (hyperhidrotic) hands and feet.

Frequency

The prevalence of Camurati-Engelmann disease is unknown. More than 300 cases have been reported worldwide.

Causes

Mutations in the TGFB1 gene cause Camurati-Engelmann disease. The TGFB1 gene provides instructions for producing a protein called transforming growth factor beta-1 (TGFβ-1). The TGFβ-1 protein triggers chemical signals that regulate various cell activities, including the growth and division (proliferation) of cells, the maturation of cells
to carry out specific functions (differentiation), cell movement (motility), and controlled cell death (apoptosis).

The TGFβ-1 protein is found throughout the body but is particularly abundant in tissues that make up the skeleton, where it helps regulate the formation and growth of bone and cartilage, a tough, flexible tissue that makes up much of the skeleton during early development. TGFβ-1 is involved in different processes in other tissues.

The TGFβ-1 protein is a protein that is found throughout the body but is particularly abundant in tissues that make up the skeleton, where it helps regulate the formation and growth of bone and cartilage, a tough, flexible tissue that makes up much of the skeleton during early development. TGFβ-1 is involved in different processes in other tissues.

The TGFβ-1 protein is found throughout the body but is particularly abundant in tissues that make up the skeleton, where it helps regulate the formation and growth of bone and cartilage, a tough, flexible tissue that makes up much of the skeleton during early development. TGFβ-1 is involved in different processes in other tissues.

The TGFB1 gene mutations that cause Camurati-Engelmann disease result in the production of an overly active TGFβ-1 protein. This abnormal TGFβ-1 protein activity causes an increase in signaling, which leads to more bone formation. As a result, the bones in the arms, legs, and skull are thicker than normal, contributing to the movement and neurological problems often experienced by individuals with Camurati-Engelmann disease.

Some individuals with Camurati-Engelmann disease do not have an identified mutation in the TGFB1 gene. In these cases, the cause of the condition is unknown.

**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.

In some cases, an affected person inherits the mutation from one affected parent. Other cases result from new mutations in the gene and occur in people with no history of the disorder in their family.

Some people who have the altered gene never develop the condition, a situation known as reduced penetrance.

**Other Names for This Condition**

- Camurati-Engelmann syndrome
- CED
- diaphyseal dysplasia
- diaphyseal hyperostosis
- diaphyseal osteosclerosis
- Engelmann disease
- PDD
- progressive diaphyseal dysplasia
Diagnosis & Management

Genetic Testing Information

• What is genetic testing?
  https://primer/testing/genetictesting

• Genetic Testing Registry: Diaphyseal dysplasia

Research Studies from ClinicalTrials.gov

• ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22Camurati-Engelmann+disease%22+OR+%22diaphyseal+dysplasia%22

Other Diagnosis and Management Resources

• GeneReview: Camurati-Engelmann Disease
  https://www.ncbi.nlm.nih.gov/books/NBK1156

• MedlinePlus Encyclopedia: Bone Mineral Density Test
  https://medlineplus.gov/ency/article/007197.htm

Additional Information & Resources

Health Information from MedlinePlus

• Encyclopedia: Bone Mineral Density Test
  https://medlineplus.gov/ency/article/007197.htm

• Encyclopedia: Bone Pain or Tenderness
  https://medlineplus.gov/ency/article/003180.htm

• Health Topic: Bone Density
  https://medlineplus.gov/bonedensity.html

• Health Topic: Bone Diseases
  https://medlineplus.gov/bonediseases.html

Genetic and Rare Diseases Information Center

• Camurati-Engelmann disease
  https://rarediseases.info.nih.gov/diseases/1072/camurati-engelmann-disease

• Camurati Engelmann disease, type 2
Educational Resources

- Children's Hospital of Philadelphia: Skeletal Dysplasias
  https://www.chop.edu/conditions-diseases/skeletal-dysplasias
- KidsHealth from Nemours: Your Bones
- MalaCards: camurati-engelmann disease
  https://www.malacards.org/card/camurati_engelmann_disease
- Orphanet: Camurati-Engelmann disease
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=1328
- UCLA International Skeletal Dysplasia Registry
  https://www.uclahealth.org/ortho/isdr

Patient Support and Advocacy Resources

- National Organization for Rare Disorders (NORD)
  https://rarediseases.org/rare-diseases/camurati-engelmann-disease/

Clinical Information from GeneReviews

- Camurati-Engelmann Disease
  https://www.ncbi.nlm.nih.gov/books/NBK1156

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28Progressive+Diaphyseal+Dysplasia%5BTIAB%5D%29+OR+%28Camurati-Engelmann+disease%5BTIAB%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- CAMURATI-ENGELMANN DISEASE
  http://omim.org/entry/131300
- CAMURATI-ENGELMANN DISEASE, TYPE 2
  http://omim.org/entry/606631
Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20566907

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15894597
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2564495/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12493741

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15326622

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/28261436
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5334020/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/22127241


Reviewed: November 2017
Published: January 29, 2019

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