Costello syndrome

Costello syndrome is a disorder that affects many parts of the body. This condition is characterized by delayed development and intellectual disability, loose folds of skin (which are especially noticeable on the hands and feet), unusually flexible joints, and distinctive facial features including a large mouth with full lips. Heart problems are common, including an abnormal heartbeat (arrhythmia), structural heart defects, and a type of heart disease that enlarges and weakens the heart muscle (hypertrophic cardiomyopathy). Infants with Costello syndrome may be larger than average at birth, but most have difficulty feeding and grow more slowly than other children. People with this condition have relatively short stature and may have reduced growth hormone levels. Other signs and symptoms of Costello syndrome can include tight Achilles tendons (which connect the calf muscles to the heel), weak muscle tone (hypotonia), a structural abnormality of the brain called a Chiari I malformation, skeletal abnormalities, dental problems, and problems with vision.

Beginning in early childhood, people with Costello syndrome are at an increased risk of developing certain cancerous and noncancerous tumors. The most common noncancerous tumors associated with this condition are papillomas, which are small, wart-like growths that usually develop around the nose and mouth or near the anus. The most common cancerous tumor associated with Costello syndrome is a childhood cancer called rhabdomyosarcoma, which begins in muscle tissue. Neuroblastoma, a tumor that arises in developing nerve cells, also has been reported in children and adolescents with this syndrome. In addition, some teenagers with Costello syndrome have developed transitional cell carcinoma, a form of bladder cancer that is usually seen in older adults.

The signs and symptoms of Costello syndrome overlap significantly with those of two other genetic conditions, cardiofaciocutaneous syndrome (CFC syndrome) and Noonan syndrome. In affected infants, it can be difficult to tell the three conditions apart based on their physical features. However, the conditions can be distinguished by their genetic cause and by specific patterns of signs and symptoms that develop later in childhood.

Frequency

This condition is very rare; it probably affects 200 to 300 people worldwide. Reported estimates of Costello syndrome prevalence range from 1 in 300,000 to 1 in 1.25 million people.

Causes

Mutations in the HRAS gene cause Costello syndrome. This gene provides instructions for making a protein called H-Ras, which is part of a pathway that helps control cell
growth and division. Mutations that cause Costello syndrome lead to the production of an H-Ras protein that is abnormally turned on (active). The overactive protein directs cells to grow and divide constantly, which can lead to the development of cancerous and noncancerous tumors. It is unclear how mutations in the HRAS gene cause the other features of Costello syndrome, but many of the signs and symptoms probably result from cell overgrowth and abnormal cell division.

Some people with signs and symptoms like those of Costello syndrome do not have an identified mutation in the HRAS gene. These individuals may actually have CFC syndrome or Noonan syndrome, which are caused by mutations in related genes. The proteins produced from these genes interact with one another and with the H-Ras protein as part of the same cell growth and division pathway. These interactions help explain why mutations in different genes can cause conditions with overlapping signs and symptoms.

Inheritance Pattern

Costello syndrome is considered to be an autosomal dominant condition, which means one copy of the altered gene in each cell is sufficient to cause the disorder. Almost all reported cases have resulted from new gene mutations and have occurred in people with no history of the disorder in their family.

Other Names for This Condition

• faciocutaneoskeletal syndrome
• FCS syndrome

Diagnosis & Management

Genetic Testing Information

• What is genetic testing?
  /primer/testing/genetictesting
• Genetic Testing Registry: Costello syndrome

Research Studies from ClinicalTrials.gov

• ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22costello+syndrome%22

Other Diagnosis and Management Resources

• GeneReview: Costello Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1507
Additional Information & Resources

Health Information from MedlinePlus

- Health Topic: Cancer
  https://medlineplus.gov/cancer.html
- Health Topic: Congenital Heart Defects
  https://medlineplus.gov/congenitalheartdefects.html
- Health Topic: Developmental Disabilities
  https://medlineplus.gov/developmentaldisabilities.html
- Health Topic: Skin Conditions
  https://medlineplus.gov/skinconditions.html

Genetic and Rare Diseases Information Center

- Costello syndrome

Educational Resources

- MalaCards: costello syndrome
  https://www.malacards.org/card/costello_syndrome
- Orphanet: Costello syndrome
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=3071

Patient Support and Advocacy Resources

- American Heart Association
  https://www.heart.org/en/health-topics/congenital-heart-defects
- Children's Craniofacial Association
  https://ccakids.org/
- Costello Syndrome Family Network
  http://costellosyndromeusa.com/
- National Organization for Rare Disorders (NORD)
  https://rarediseases.org/rare-diseases/costello-syndrome/
- RASopathiesNet
  https://rasopathiesnet.org/
- Resource list from the University of Kansas Medical Center
  http://www.kumc.edu/gec/support/costello.html
- The MAGIC Foundation
  https://www.magicfoundation.org/
Clinical Information from GeneReviews

• Costello Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1507

Scientific Articles on PubMed

• PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28costello+syndrome%5BTIAB%5D%29+AND+english%5BLa%5D+AND+human%5Bmh%5D+AND+%22last+1800+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

• COSTELLO SYNDROME
  http://omim.org/entry/218040

Sources for This Summary

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