Tetra-amelia syndrome

Tetra-amelia syndrome is a very rare disorder characterized by the absence of all four limbs. ("Tetra" is the Greek word for "four," and "amelia" refers to the failure of an arm or leg to develop before birth.) This syndrome can also cause severe malformations of other parts of the body, including the face and head, heart, nervous system, skeleton, and genitalia. The lungs are underdeveloped in many cases, which makes breathing difficult or impossible. Because children with tetra-amelia syndrome have such serious medical problems, most are stillborn or die shortly after birth.

Frequency

Tetra-amelia syndrome has been reported in only a few families worldwide.

Genetic Changes

Researchers have found a mutation in the \textit{WNT3} gene in people with tetra-amelia syndrome from one large family. This gene is part of a family of WNT genes that play critical roles in development before birth. The protein produced from the \textit{WNT3} gene is involved in the formation of the limbs and other body systems during embryonic development. Mutations in the \textit{WNT3} gene prevent cells from producing functional WNT3 protein, which disrupts normal limb formation and leads to the other serious birth defects associated with tetra-amelia syndrome.

In other affected families, the cause of tetra-amelia syndrome has not been determined. Researchers believe that unidentified mutations in \textit{WNT3} or other genes involved in limb development are probably responsible for the disorder in these cases.

Inheritance Pattern

In most of the families reported so far, tetra-amelia syndrome appears to have an autosomal recessive pattern of inheritance. Autosomal recessive inheritance means both copies of the gene in each cell have mutations. The parents of an individual with tetra-amelia syndrome each carry one copy of the mutated gene, but do not show signs and symptoms of the condition.

Other Names for This Condition

- Tetra-amelia
- Tetra-amelia, autosomal recessive
Diagnosis & Management

Genetic Testing

- Genetic Testing Registry: Tetraamelia, autosomal recessive

Other Diagnosis and Management Resources

- GeneReview: Tetra-Amelia Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1276

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

MedlinePlus

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

Genetic and Rare Diseases Information Center

- Tetra-amelia syndrome

Educational Resources

- Disease InfoSearch: Tetraamelia autosomal recessive
  http://www.diseaseinfosearch.org/Tetraamelia+autosomal+recessive/7034

- Lucile Packard Children’s Hospital: Congenital Limb Defects

- Orphanet: Tetraamelia-multiple malformations syndrome
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=3301
Patient Support and Advocacy Resources

- Resource list from the University of Kansas Medical Center: Limb Anomalies
  http://www.kumc.edu/gec/support/limb.html
- The Compassionate Friends
  https://www.compassionatefriends.org/

GeneReviews

- Tetra-Amelia Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1276

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28tetra-amelia%5BTIAB%5D %29+OR+%28tetraamelia%5BTIAB%5D%29%29+AND+english%5Bla%5D+AND
  +human%5Bmhm%5D

OMIM

- TETRAAMELIA SYNDROME, AUTOSOMAL RECESSIVE
  http://omim.org/entry/273395

MedGen

- Tetra-amelia with pulmonary hypoplasia

Sources for This Summary

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Reprinted from Genetics Home Reference:

Reviewed: February 2008
Published: May 29, 2018

Lister Hill National Center for Biomedical Communications
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