

Achondrogenesis

Description

Achondrogenesis is a group of severe disorders that affect cartilage and bone development. These conditions are characterized by a small body, short limbs, and other skeletal abnormalities. As a result of serious health problems, infants with achondrogenesis usually die before birth, are stillborn, or die soon after birth from respiratory failure. However, some infants have lived for a short time with intensive medical support.

Researchers have described at least three forms of achondrogenesis, designated as type 1A, type 1B, and type 2. The types are distinguished by their signs and symptoms, inheritance pattern, and genetic cause. However, types 1A and 1B are often hard to tell apart without genetic testing.

Achondrogenesis type 1A, which is also called the Houston-Harris type, is the least well understood of the three forms. Affected infants have extremely short limbs, a narrow chest, short ribs that fracture easily, and a lack of normal bone formation (ossification) in the skull, spine, and pelvis.

Achondrogenesis type 1B, also known as the Parenti-Fraccaro type, is characterized by extremely short limbs, a narrow chest, and a prominent, rounded abdomen. The fingers and toes are short and the feet may turn inward and upward (clubfeet). Affected infants frequently have a soft out-pouching around the belly-button (an umbilical hernia) or near the groin (an inguinal hernia).

Infants with achondrogenesis type 2, which is sometimes called the Langer-Saldino type, have short arms and legs, a narrow chest with short ribs, and underdeveloped lungs. This condition is also associated with a lack of ossification in the spine and pelvis. Distinctive facial features include a prominent forehead, a small chin, and, in some cases, an opening in the roof of the mouth (a cleft palate). The abdomen is enlarged, and affected infants often have a condition called hydrops fetalis, in which excess fluid builds up in the body before birth.

Frequency

Achondrogenesis types 1A and 1B are rare genetic disorders; their incidence is unknown. Combined, achondrogenesis type 2 and hypochondrogenesis (a similar skeletal disorder) occur in 1 in 40,000 to 60,000 newborns.

Causes

Mutations in the *TRIP11*, *SLC26A2*, and *COL2A1* genes cause achondrogenesis type 1A, type 1B, and type 2, respectively.

The genetic cause of achondrogenesis type 1A was unknown until recently, when researchers discovered that the condition can result from mutations in the *TRIP11* gene. This gene provides instructions for making a protein called GMAP-210. This protein plays a critical role in the Golgi apparatus, a cell structure in which newly produced proteins are modified so they can carry out their functions. Mutations in the *TRIP11* gene prevent the production of functional GMAP-210, which alters the structure and function of the Golgi apparatus. Researchers suspect that cells called chondrocytes in the developing skeleton may be most sensitive to these changes. Chondrocytes give rise to cartilage, 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. Malfunction of the Golgi apparatus in chondrocytes likely underlies the problems with bone formation in achondrogenesis type 1A.

Achondrogenesis type 1B is the most severe of a spectrum of 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. Mutations in the *SLC26A2* gene cause the skeletal problems characteristic of achondrogenesis type 1B by disrupting the structure of developing cartilage, which prevents bones from forming properly.

Achondrogenesis type 2 is one of several skeletal disorders that result from mutations in the *COL2A1* gene. This gene provides instructions for making a protein that forms type II collagen. This type of collagen is found mostly in cartilage and in the clear gel that fills the eyeball (the vitreous). It is essential for the normal development of bones and other connective tissues that form the body's supportive framework. Mutations in the *COL2A1* gene interfere with the assembly of type II collagen molecules, which prevents bones and other connective tissues from developing properly.

[Learn more about the genes associated with Achondrogenesis](#)

- COL2A1
- SLC26A2
- TRIP11

Inheritance

Achondrogenesis type 1A and type 1B both have an autosomal recessive pattern of inheritance, which means both copies of the *TRIP11* or *SLC26A2* gene in each cell have mutations. Most often, the parents of an individual with an autosomal recessive condition each carry one copy of the mutated gene but do not show signs and symptoms of the condition.

Achondrogenesis type 2 is considered an autosomal dominant disorder because one copy of the altered gene in each cell is sufficient to cause the condition. It is almost always caused by new mutations in the *COL2A1* gene and typically occurs in people with no history of the disorder in their family.

Other Names for This Condition

- Achondrogenesis syndrome

Additional Information & Resources

Genetic Testing Information

- Genetic Testing Registry: Achondrogenesis type II (<https://www.ncbi.nlm.nih.gov/gtr/conditions/C0220685/>)
- Genetic Testing Registry: Achondrogenesis, type IA (<https://www.ncbi.nlm.nih.gov/gtr/conditions/C0265273/>)
- Genetic Testing Registry: Achondrogenesis, type IB (<https://www.ncbi.nlm.nih.gov/gtr/conditions/C0265274/>)

Genetic and Rare Diseases Information Center

- Achondrogenesis (<https://rarediseases.info.nih.gov/diseases/2882/index>)

Patient Support and Advocacy Resources

- National Organization for Rare Disorders (NORD) (<https://rarediseases.org/>)

Catalog of Genes and Diseases from OMIM

- ACHONDROGENESIS, TYPE IA; ACG1A (<https://omim.org/entry/200600>)
- ACHONDROGENESIS, TYPE II; ACG2 (<https://omim.org/entry/200610>)
- ACHONDROGENESIS, TYPE IB; ACG1B (<https://omim.org/entry/600972>)

Scientific Articles on PubMed

- PubMed (<https://pubmed.ncbi.nlm.nih.gov/?term=%28achondrogenesis%5BTIAB%5D%29+AND+english%5BIa%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D>)

References

- Faivre L, Le Merrer M, Douvier S, Laurent N, Thauvin-Robinet C, Rousseau T, Vereecke I, Sagot P, Delezoide AL, Coucke P, Mortier G. Recurrence of achondrogenesis type II within the same family: evidence for germline mosaicism. *Am J Med Genet A*. 2004 Apr 30;126A(3):308-12. doi: 10.1002/ajmg.a.20597. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/15054848>)
- Grigeliuniene G, Geiger S, Papadogiannakis N, Makitie O, Nishimura G, Nordgren A, Conner P. The phenotype range of achondrogenesis 1A. *Am J Med Genet A*. 2013 Oct;161A(10):2554-8. doi: 10.1002/ajmg.a.36106. Epub 2013 Aug 16. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/23956106>)
- Kapur RP. Achondrogenesis. *Pediatr Dev Pathol*. 2007 Jul-Aug;10(4):253-5. doi:10.2350/07-01-0216.1. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/17638434>)
- Korkko J, Cohn DH, Ala-Kokko L, Krakow D, Prockop DJ. Widely distributed mutations in the COL2A1 gene produce achondrogenesis type II/hypochondrogenesis. *Am J Med Genet*. 2000 May 15;92(2):95-100. doi:10.1002/(sici)1096-8628(20000515)92:23.0.co;2-9. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/10797431>)
- Rossi A, Superti-Furga A. Mutations in the diastrophic dysplasia sulfate transporter (DTDST) gene (SLC26A2): 22 novel mutations, mutation review, associated skeletal phenotypes, and diagnostic relevance. *Hum Mutat*. 2001 Mar;17(3):159-71. doi: 10.1002/humu.1. Erratum In: *Hum Mutat* 2001;18(1):82. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/11241838>)
- Smits P, Bolton AD, Funari V, Hong M, Boyden ED, Lu L, Manning DK, Dwyer ND, Moran JL, Prysak M, Merriman B, Nelson SF, Bonafe L, Superti-Furga A, Ikegawa S, Krakow D, Cohn DH, Kirchhausen T, Warman ML, Beier DR. Lethal skeletal dysplasia in mice and humans lacking the golgin GMAP-210. *N Engl J Med*. 2010 Jan 21;362(3):206-16. doi: 10.1056/NEJMoa0900158. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/20089971>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3108191/>)
- Superti-Furga A, Hastbacka J, Wilcox WR, Cohn DH, van der Harten HJ, Rossi A, Blau N, Rimoin DL, Steinmann B, Lander ES, Gitzelmann R. Achondrogenesis type IB is caused by mutations in the diastrophic dysplasia sulphate transporter gene. *Nat Genet*. 1996 Jan;12(1):100-2. doi: 10.1038/ng0196-100. No abstract available. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/8528239>)

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