



Carpenter syndrome

Carpenter syndrome is a condition characterized by the premature fusion of certain skull bones (craniosynostosis), abnormalities of the fingers and toes, and other developmental problems.

Craniosynostosis prevents the skull from growing normally, frequently giving the head a pointed appearance (acrocephaly). In severely affected individuals, the abnormal fusion of the skull bones results in a deformity called a cloverleaf skull. Craniosynostosis can cause differences between the two sides of the head and face (craniofacial asymmetry). Early fusion of the skull bones can affect the development of the brain and lead to increased pressure within the skull (intracranial pressure). Premature fusion of the skull bones can cause several characteristic facial features in people with Carpenter syndrome. Distinctive facial features may include a flat nasal bridge, outside corners of the eyes that point downward (down-slanting palpebral fissures), low-set and abnormally shaped ears, underdeveloped upper and lower jaws, and abnormal eye shape. Some affected individuals also have dental abnormalities including small primary (baby) teeth. Vision problems also frequently occur.

Abnormalities of the fingers and toes include fusion of the skin between two or more fingers or toes (cutaneous syndactyly), unusually short fingers or toes (brachydactyly), or extra fingers or toes (polydactyly). In Carpenter syndrome, cutaneous syndactyly is most common between the third (middle) and fourth (ring) fingers, and polydactyly frequently occurs next to the big or second toe or the fifth (pinky) finger.

People with Carpenter syndrome often have intellectual disability, which can range from mild to profound. However, some individuals with this condition have normal intelligence. The cause of intellectual disability is unknown, as the severity of craniosynostosis does not appear to be related to the severity of intellectual disability.

Other features of Carpenter syndrome include obesity that begins in childhood, a soft out-pouching around the belly-button (umbilical hernia), hearing loss, and heart defects. Additional skeletal abnormalities such as deformed hips, a rounded upper back that also curves to the side (kyphoscoliosis), and knees that are angled inward (genu valgum) frequently occur. Nearly all affected males have genital abnormalities, most frequently undescended testes (cryptorchidism).

A few people with Carpenter syndrome have organs or tissues within their chest and abdomen that are in mirror-image reversed positions. This abnormal placement may affect several internal organs (situs inversus); just the heart (dextrocardia), placing the heart on the right side of the body instead of on the left; or only the major (great) arteries of the heart, altering blood flow.

The signs and symptoms of this disorder vary considerably, even within the same family. The life expectancy for individuals with Carpenter syndrome is shortened but extremely variable.

The signs and symptoms of Carpenter syndrome are similar to another genetic condition called Greig cephalopolysyndactyly syndrome. The overlapping features, which include craniosynostosis, polydactyly, and heart abnormalities, can cause these two conditions to be misdiagnosed; genetic testing is often required for an accurate diagnosis.

Frequency

Carpenter syndrome is thought to be a rare condition; approximately 70 cases have been described in the scientific literature.

Causes

Mutations in the *RAB23* or *MEGF8* gene cause Carpenter syndrome.

The *RAB23* gene provides instructions for making a protein that is involved in a process called vesicle trafficking, which moves proteins and other molecules within cells in sac-like structures called vesicles. The Rab23 protein transports vesicles from the cell membrane to their proper location inside the cell. Vesicle trafficking is important for the transport of materials that are needed to trigger signaling during development. For example, the Rab23 protein regulates a developmental pathway called the hedgehog signaling pathway that is critical in cell growth (proliferation), cell specialization, and the normal shaping (patterning) of many parts of the body.

The *MEGF8* gene provides instructions for making a protein whose function is unclear. Based on its structure, the Megf8 protein may be involved in cell processes such as sticking cells together (cell adhesion) and helping proteins interact with each other. Researchers also suspect that the Megf8 protein plays a role in normal body patterning.

Mutations in the *RAB23* or *MEGF8* gene lead to the production of proteins with little or no function. It is unclear how disruptions in protein function lead to the features of Carpenter syndrome, but it is likely that interference with normal body patterning plays a role. For reasons that are unknown, people with *MEGF8* gene mutations are more likely to have dextrocardia and other organ positioning abnormalities and less severe craniosynostosis than individuals with *RAB23* gene mutations.

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

- ACPS II
- acrocephalopolysyndactyly 2
- acrocephalopolysyndactyly type II
- acrocephalosyndactyly, type II
- type II acrocephalosyndactyly

Diagnosis & Management

Genetic Testing Information

- What is genetic testing?
[/primer/testing/genetic-testing](#)
- Genetic Testing Registry: Carpenter syndrome
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C1275078/>
- Genetic Testing Registry: Carpenter syndrome 2
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C3554247/>

Research Studies from ClinicalTrials.gov

- ClinicalTrials.gov
<https://clinicaltrials.gov/ct2/results?cond=%22Carpenter+syndrome%22+OR+%22acrocephalopolysyndactyly+type+II%22+OR+%22Acrocephalosyndactyly%22>

Other Diagnosis and Management Resources

- MedlinePlus Encyclopedia: Craniosynostosis Repair
<https://medlineplus.gov/ency/article/007364.htm>
- MedlinePlus Encyclopedia: Dextrocardia
<https://medlineplus.gov/ency/article/007326.htm>

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Craniosynostosis
<https://medlineplus.gov/ency/article/001590.htm>
- Encyclopedia: Craniosynostosis Repair
<https://medlineplus.gov/ency/article/007364.htm>
- Encyclopedia: Dextrocardia
<https://medlineplus.gov/ency/article/007326.htm>
- Encyclopedia: Skull of a Newborn (image)
<https://medlineplus.gov/ency/imagepages/1127.htm>

- Encyclopedia: Umbilical Hernia
<https://medlineplus.gov/ency/article/000987.htm>
- Encyclopedia: Webbing of the Fingers or Toes
<https://medlineplus.gov/ency/article/003289.htm>
- Health Topic: Craniofacial Abnormalities
<https://medlineplus.gov/craniofacialabnormalities.html>

Genetic and Rare Diseases Information Center

- Carpenter syndrome
<https://rarediseases.info.nih.gov/diseases/6003/carpenter-syndrome>

Additional NIH Resources

- National Institute of Neurological Disorders and Stroke: Craniosynostosis Information Page
<https://www.ninds.nih.gov/Disorders/All-Disorders/Craniosynostosis-Information-Page>

Educational Resources

- American Society for Surgery of the Hand: Congenital Hand Differences
<http://handcare.assh.org/Anatomy/Details-Page/articleId/39392>
- Boston Children's Hospital: Craniosynostosis
<http://www.childrenshospital.org/conditions-and-treatments/conditions/c/craniosynostosis>
- Boston Children's Hospital: Polydactyly
<http://www.childrenshospital.org/conditions-and-treatments/conditions/p/polydactyly>
- Centers for Disease Control and Prevention: Congenital Heart Defects
<https://www.cdc.gov/ncbddd/heartdefects/>
- Centers for Disease Control and Prevention: Facts about dextro-Transposition of the Great Arteries (d-TGA)
<https://www.cdc.gov/ncbddd/heartdefects/d-tga.html>
- Cincinnati Children's Hospital: Craniosynostosis
<https://www.cincinnatichildrens.org/health/c/craniosynostosis>
- Kennedy Krieger Institute: Intellectual Disability
<https://www.kennedykrieger.org/patient-care/conditions/intellectual-disability>
- MalaCards: carpenter syndrome 1
https://www.malacards.org/card/carpenter_syndrome_1
- MalaCards: carpenter syndrome 2
https://www.malacards.org/card/carpenter_syndrome_2

- Orphanet: Carpenter syndrome
https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=65759
- Seattle Children's Hospital: Syndactyly
<https://www.seattlechildrens.org/conditions/bone-joint-muscle-conditions/syndactyly>

Patient Support and Advocacy Resources

- AboutFace
<https://www.aboutface.ca/>
- American Association on Intellectual and Developmental Disabilities (AAIDD)
<https://www.aaid.org/>
- American Heart Association
<https://www.heart.org/>
- Children's Craniofacial Association
<https://ccakids.org/syndromes.html>
- Children's Craniofacial Association: Craniosynostosis
<https://ccakids.org/craniosynostosis.html>
- National Organization for Rare Disorders (NORD)
<https://rarediseases.org/rare-diseases/carpenter-syndrome/>
- Resource List from the University of Kansas Medical Center: Facial Anomalies/
Craniofacial Conditions
<http://www.kumc.edu/gec/support/craniofa.html>

Scientific Articles on PubMed

- PubMed
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28carpenter+syndrome%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

- CARPENTER SYNDROME 1
<http://omim.org/entry/201000>
- CARPENTER SYNDROME 2
<http://omim.org/entry/614976>

Medical Genetics Database from MedGen

- Carpenter syndrome
<https://www.ncbi.nlm.nih.gov/medgen/226897>
- Carpenter syndrome 2
<https://www.ncbi.nlm.nih.gov/medgen/767161>

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