Cornelia de Lange syndrome

Cornelia de Lange syndrome is a developmental disorder that affects many parts of the body. The features of this disorder vary widely among affected individuals and range from relatively mild to severe.

Cornelia de Lange syndrome is characterized by slow growth before and after birth leading to short stature; intellectual disability that is usually moderate to severe; and abnormalities of bones in the arms, hands, and fingers. Most people with Cornelia de Lange syndrome also have distinctive facial features, including arched eyebrows that often meet in the middle (synophrys), long eyelashes, low-set ears, small and widely spaced teeth, and a small and upturned nose. Many affected individuals also have behavior problems similar to autism, a developmental condition that affects communication and social interaction.

Additional signs and symptoms of Cornelia de Lange syndrome can include excessive body hair (hypertrichosis), an unusually small head (microcephaly), hearing loss, and problems with the digestive tract. Some people with this condition are born with an opening in the roof of the mouth called a cleft palate. Seizures, heart defects, and eye problems have also been reported in people with this condition.

Frequency

Although the exact incidence is unknown, Cornelia de Lange syndrome likely affects 1 in 10,000 to 30,000 newborns. The condition is probably underdiagnosed because affected individuals with mild or uncommon features may never be recognized as having Cornelia de Lange syndrome.

Causes

Cornelia de Lange syndrome can result from mutations in at least five genes: NIPBL, SMC1A, HDAC8, RAD21, and SMC3. Mutations in the NIPBL gene have been identified in more than half of all people with this condition; mutations in the other genes are much less common.

The proteins produced from all five genes contribute to the structure or function of the cohesin complex, a group of proteins with an important role in directing development before birth. Within cells, the cohesin complex helps regulate the structure and organization of chromosomes, stabilize cells' genetic information, and repair damaged DNA. The cohesin complex also regulates the activity of certain genes that guide the development of limbs, face, and other parts of the body.
Mutations in the NIPBL, SMC1A, HDAC8, RAD21, and SMC3 genes cause Cornelia de Lange syndrome by impairing the function of the cohesin complex, which disrupts gene regulation during critical stages of early development.

The features of Cornelia de Lange syndrome vary widely, and the severity of the disorder can differ even in individuals with the same gene mutation. Researchers suspect that additional genetic or environmental factors may be important for determining the specific signs and symptoms in each individual. In general, SMC1A, RAD21, and SMC3 gene mutations cause milder signs and symptoms than NIPBL gene mutations. Mutations in the HDAC8 gene cause a somewhat different set of features, including delayed closure of the "soft spot" on the head (the anterior fontanelle) in infancy, widely spaced eyes, and dental abnormalities. Like affected individuals with NIPBL gene mutations, those with HDAC8 gene mutations may have significant intellectual disability.

In about 30 percent of cases, the cause of Cornelia de Lange syndrome is unknown. Researchers are looking for additional changes in the five known genes, as well as mutations in other genes, that may cause this condition.

Inheritance Pattern

When Cornelia de Lange syndrome is caused by mutations in the NIPBL, RAD21, or SMC3 gene, the condition is considered to have an autosomal dominant pattern of inheritance. Autosomal dominant inheritance means one copy of the altered gene in each cell is sufficient to cause the disorder. Most cases result from new gene mutations and occur in people with no history of the condition in their family.

When Cornelia de Lange syndrome is caused by mutations in the HDAC8 or SMC1A gene, the condition has an X-linked dominant pattern of inheritance. A condition is considered X-linked if the mutated gene that causes the disorder is located on the X chromosome, one of the two sex chromosomes. Studies of X-linked Cornelia de Lange syndrome indicate that one copy of the altered gene in each cell may be sufficient to cause the condition. Unlike X-linked recessive conditions, in which males are more frequently affected or experience more severe symptoms than females, X-linked dominant Cornelia de Lange syndrome appears to affect males and females similarly. Most cases result from new mutations in the HDAC8 or SMC1A gene and occur in people with no history of the condition in their family.

Other Names for This Condition

- BDLS
- Brachmann-de Lange syndrome
- CdLS
- de Lange syndrome
- typus degenerativus amstelodamensis
Diagnosis & Management

Genetic Testing Information

- What is genetic testing?
  /primer/testing/genetictesting

- Genetic Testing Registry: Cornelia de Lange Syndrome

- Genetic Testing Registry: De Lange syndrome

Research Studies from ClinicalTrials.gov

- ClinicalTrials.gov
  https://clinicaltrials.gov/ct2/results?cond=%22Cornelia+de+Lange+syndrome%22

Other Diagnosis and Management Resources

- CdLS World: CdLS Diagnostic Criteria Checklist
  https://www.cdlsworld.org/xwiki/bin/view/CdLSWorld/topicDiagnosisCriteria

- CdLS World: Management and Treatment Guidelines for CdLS
  https://www.cdlsworld.org/xwiki/bin/view/CdLSWorld/TreatmentGuidelinesPreventiveCare

- CdLS World: Treatment Protocols
  https://www.cdlsworld.org/xwiki/bin/view/CdLSWorld/CdLSTreatmentProtocols

- Cornelia de Lange Syndrome Foundation: Diagnosis of CdLS
  https://www.cdlsusa.org/diagnosis-of-cdls/

- Cornelia de Lange Syndrome Foundation: Treatment Protocols
  https://www.cdlsusa.org/treatment-protocols-2/

- GeneReview: Cornelia de Lange Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1104

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Autism
  https://medlineplus.gov/ency/article/001526.htm

- Encyclopedia: Microcephaly
  https://medlineplus.gov/ency/article/003272.htm

- Health Topic: Developmental Disabilities
  https://medlineplus.gov/developmentaldisabilities.html
Genetic and Rare Diseases Information Center

- Cornelia de Lange syndrome
  https://rarediseases.info.nih.gov/diseases/10109/cornelia-de-lange-syndrome

Educational Resources

- MalaCards: cornelia de lange syndrome 1
  https://www.malacards.org/card/cornelia_de_lange_syndrome_1
- Orphanet: Cornelia de Lange syndrome
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=199
- UC Davis Children’s Hospital

Patient Support and Advocacy Resources

- CdLS UK and Ireland: The Cornelia de Lange Syndrome Foundation
  http://www.cdls.org.uk
- CdLS World: The International Federation of National CdLS Support Organizations
  https://www.cdlsworld.org/xwiki/bin/view/Main/WebHome
- Cornelia de Lange Syndrome Foundation
  https://www.cdlsusa.org/
- National Organization for Rare Disorders (NORD)
  https://rarediseases.org/rare-diseases/cornelia-de-lange-syndrome/
- Resource list from the University of Kansas Medical Center
  http://www.kumc.edu/gec/support/cornelia.html

Clinical Information from GeneReviews

- Cornelia de Lange Syndrome
  https://www.ncbi.nlm.nih.gov/books/NBK1104

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28De+Lange+Syndrome%5BMAJR%5D%29+AND+%28Cornelia+de+Lange+syndrome%5BTIAB%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1080+days%22+AND+indent%5D

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Catalog of Genes and Diseases from OMIM

- CORNELIA DE LANGE SYNDROME 1
  http://omim.org/entry/122470

- CORNELIA DE LANGE SYNDROME 2
  http://omim.org/entry/300590

- CORNELIA DE LANGE SYNDROME 3
  http://omim.org/entry/610759

- CORNELIA DE LANGE SYNDROME 4
  http://omim.org/entry/614701

- CORNELIA DE LANGE SYNDROME 5
  http://omim.org/entry/300882

Medical Genetics Database from MedGen

- Cornelia de Lange Syndrome

- De Lange syndrome

Sources for This Summary


  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3443318/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/20301283

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/22633999
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3370273/


