Oculodentodigital dysplasia

Oculodentodigital dysplasia is a condition that affects many parts of the body, particularly the eyes (oculo-), teeth (dento-), and fingers (digital). Common features in people with this condition are small eyes (microphthalmia) and other eye abnormalities that can lead to vision loss. Affected individuals also frequently have tooth abnormalities, such as small or missing teeth, weak enamel, multiple cavities, and early tooth loss. Other common features of this condition include a thin nose and webbing of the skin (syndactyly) between the fourth and fifth fingers.

Less common features of oculodentodigital dysplasia include sparse hair growth (hypotrichosis), brittle nails, an unusual curvature of the fingers (camptodactyly), syndactyly of the toes, small head size (microcephaly), and an opening in the roof of the mouth (cleft palate). Some affected individuals experience neurological problems such as a lack of bladder or bowel control, difficulty coordinating movements (ataxia), abnormal muscle stiffness (spasticity), hearing loss, and impaired speech (dysarthria). A few people with oculodentodigital dysplasia also have a skin condition called palmoplantar keratoderma. Palmoplantar keratoderma causes the skin on the palms and the soles of the feet to become thick, scaly, and calloused.

Some features of oculodentodigital dysplasia are evident at birth, while others become apparent with age.

Frequency

The exact incidence of oculodentodigital dysplasia is unknown. It has been diagnosed in fewer than 1,000 people worldwide. More cases are likely undiagnosed.

Causes

Mutations in the GJA1 gene cause oculodentodigital dysplasia. The GJA1 gene provides instructions for making a protein called connexin 43. This protein forms one part (a subunit) of channels called gap junctions, which allow direct communication between cells. Gap junctions formed by connexin 43 proteins are found in many tissues throughout the body.

GJA1 gene mutations result in abnormal connexin 43 proteins. Channels formed with abnormal proteins are often permanently closed. Some mutations prevent connexin 43 proteins from traveling to the cell surface where they are needed to form channels between cells. Impaired functioning of these channels disrupts cell-to-cell communication, which likely interferes with normal cell growth and cell specialization, processes that determine the shape and function of many different parts of the body. These developmental problems cause the signs and symptoms of oculodentodigital dysplasia.
Inheritance Pattern

Most cases of oculodentodigital dysplasia are inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to cause the disorder. In some cases, an affected person inherits the mutation from one affected parent. Other cases result from new mutations in the gene and occur in people with no history of the disorder in their family.

Less commonly, oculodentodigital dysplasia can be 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. Fewer than ten cases of autosomal recessive oculodentodigital dysplasia have been reported.

Other Names for This Condition

• oculo-dento-digital dysplasia
• oculo-dento-osseous dysplasia
• oculodentodigital syndrome
• oculodentoosseous dysplasia
• ODD syndrome
• ODDD
• ODOD
• osseous-oculo-dental dysplasia

Diagnosis & Management

Genetic Testing

• Genetic Testing Registry: Oculodentodigital dysplasia

Other Diagnosis and Management Resources

• MedlinePlus Encyclopedia: Webbing of the fingers or toes
  https://medlineplus.gov/ency/article/003289.htm
• UC Davis Children's Hospital: Cleft and Craniofacial Reconstruction
  https://www.ucdmc.ucdavis.edu/children/clincial_services/cleft_craniofacial/
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

Health Information from MedlinePlus

• Encyclopedia: Webbing of the fingers or toes
  https://medlineplus.gov/ency/article/003289.htm

• Health Topic: Cleft Lip and Palate
  https://medlineplus.gov/cleftlipandpalate.html

• Health Topic: Tooth Disorders
  https://medlineplus.gov/toothdisorders.html

Genetic and Rare Diseases Information Center

• Oculodentodigital dysplasia
  https://rarediseases.info.nih.gov/diseases/7239/oculodentodigital-dysplasia

Educational Resources

• Boston Children's Hospital: Syndactyly
  http://www.childrenshospital.org/conditions-and-treatments/conditions/s/syndactyly

• MalaCards: oculodentodigital dysplasia
  http://www.malacards.org/card/oculodentodigital_dysplasia

• March of Dimes: Cleft Lip and Cleft Palate

• Orphanet: Oculodentodigital dysplasia
  https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=2710

• Washington University, St. Louis: Neuromuscular Disease Center
  https://neuromuscular.wustl.edu/spinal/fsp.html#oddd
Patient Support and Advocacy Resources

- American Cleft Palate-Craniofacial Association
  https://cleftline.org/
- Children's Craniofacial Association
  https://ccakids.org/
- National Foundation for Ectodermal Dysplasias
  https://www.nfed.org/
- National Organization for Rare Disorders (NORD)
  https://rarediseases.org/rare-diseases/oculo-dento-digital-dysplasia/
- The Foundation Fighting Blindness
  http://ffb.ca/
- University of Kansas Medical Center Resource List: Hard of Hearing/Deafness
  http://www.kumc.edu/gec/support/hearing.html

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28oculodentodigital+dysplasia+%5BTIAB%5D%29+OR+%28oculo-dento-digital+dysplasia%5BTIAB%5D%29+OR+%28oddd%5BTIAB%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1080+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- OCULODENTODIGITAL DYSPLASIA
  http://omim.org/entry/164200

Sources for This Summary

  
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16492141
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1383703/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12021949

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/12457340
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC379233/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15879313

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/17256797

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  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15551259

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