



Dentinogenesis imperfecta

Dentinogenesis imperfecta is a disorder of tooth development. This condition causes the teeth to be discolored (most often a blue-gray or yellow-brown color) and translucent. Teeth are also weaker than normal, making them prone to rapid wear, breakage, and loss. These problems can affect both primary (baby) teeth and permanent teeth.

Researchers have described three types of dentinogenesis imperfecta with similar dental abnormalities. Type I occurs in people who have osteogenesis imperfecta, a genetic condition in which bones are brittle and easily broken. Dentinogenesis imperfecta type II and type III usually occur in people without other inherited disorders. A few older individuals with type II have had progressive high-frequency hearing loss in addition to dental abnormalities, but it is not known whether this hearing loss is related to dentinogenesis imperfecta.

Some researchers believe that dentinogenesis imperfecta type II and type III, along with a condition called dentin dysplasia type II, are actually forms of a single disorder. The signs and symptoms of dentin dysplasia type II are very similar to those of dentinogenesis imperfecta. However, dentin dysplasia type II affects the primary teeth much more than the permanent teeth.

Frequency

Dentinogenesis imperfecta affects an estimated 1 in 6,000 to 8,000 people.

Causes

Mutations in the *DSPP* gene have been identified in people with dentinogenesis imperfecta type II and type III. Mutations in this gene are also responsible for dentin dysplasia type II. Dentinogenesis imperfecta type I occurs as part of osteogenesis imperfecta, which is caused by mutations in one of several other genes (most often the *COL1A1* or *COL1A2* genes).

The *DSPP* gene provides instructions for making two proteins that are essential for normal tooth development. These proteins are involved in the formation of dentin, which is a bone-like substance that makes up the protective middle layer of each tooth. *DSPP* gene mutations alter the proteins made from the gene, leading to the production of abnormally soft dentin. Teeth with defective dentin are discolored, weak, and more likely to decay and break. It is unclear whether *DSPP* gene mutations are related to the hearing loss found in a few older individuals with dentinogenesis imperfecta type II.

Inheritance Pattern

This condition is inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to cause the disorder.

Other Names for This Condition

- DGI
- hereditary opalescent dentin

Diagnosis & Management

Genetic Testing Information

- What is genetic testing?
[/primer/testing/genetic-testing](#)
- Genetic Testing Registry: Dentinogenesis imperfecta - Shield's type II
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C2973527/>
- Genetic Testing Registry: Dentinogenesis imperfecta - Shield's type III
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C0399378/>

Research Studies from ClinicalTrials.gov

- ClinicalTrials.gov
<https://clinicaltrials.gov/ct2/results?cond=%22dentinogenesis+imperfecta%22>

Other Diagnosis and Management Resources

- MedlinePlus Encyclopedia: Tooth - Abnormal Colors
<https://medlineplus.gov/ency/article/003065.htm>

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Tooth - Abnormal Colors
<https://medlineplus.gov/ency/article/003065.htm>
- Health Topic: Tooth Disorders
<https://medlineplus.gov/toothdisorders.html>

Genetic and Rare Diseases Information Center

- Dentinogenesis imperfecta
<https://rarediseases.info.nih.gov/diseases/6258/dentinogenesis-imperfecta>
- Dentinogenesis imperfecta type 2
<https://rarediseases.info.nih.gov/diseases/12796/dentinogenesis-imperfecta-type-2>
- Dentinogenesis imperfecta type 3
<https://rarediseases.info.nih.gov/diseases/10144/dentinogenesis-imperfecta-type-3>

Educational Resources

- MalaCards: dentinogenesis imperfecta
https://www.malacards.org/card/dentinogenesis_imperfecta
- Merck Manual Home Edition for Patients and Caregivers: Overview of Tooth Disorders
<https://www.merckmanuals.com/home/mouth-and-dental-disorders/tooth-disorders/overview-of-tooth-disorders>
- Orphanet: Dentin dysplasia
https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=1653
- Orphanet: Dentinogenesis imperfecta
https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=49042
- Orphanet: Dentinogenesis imperfecta type 2
https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=166260
- Orphanet: Dentinogenesis imperfecta type 3
https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=166265
- School of Dentistry, University of North Carolina at Chapel Hill
<https://www.dentistry.unc.edu/dentalprofessionals/resources/defects/di/>

Patient Support and Advocacy Resources

- National Organization for Rare Disorders (NORD): Dentinogenesis Imperfecta Type III
<https://rarediseases.org/rare-diseases/dentinogenesis-imperfecta-type-iii/>
- Osteogenesis Imperfecta Foundation
https://oif.org/wp-content/uploads/2019/08/Dental_Care.pdf

Scientific Articles on PubMed

- PubMed
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28Dentinogenesis+Imperfecta%5BMAJR%5D%29+AND+%28dentinogenesis+imperfecta%5BTIAB%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1800+days%22%5Bdp%5D>

Catalog of Genes and Diseases from OMIM

- DENTIN DYSPLASIA, TYPE II
<http://omim.org/entry/125420>
- DENTINOGENESIS IMPERFECTA 1
<http://omim.org/entry/125490>
- DENTINOGENESIS IMPERFECTA, SHIELDS TYPE III
<http://omim.org/entry/125500>

Medical Genetics Database from MedGen

- Dentinogenesis imperfecta - Shield type II
<https://www.ncbi.nlm.nih.gov/medgen/424922>

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Free article on PubMed Central: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3591212/>
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Reprinted from Genetics Home Reference:

<https://ghr.nlm.nih.gov/condition/dentinogenesis-imperfecta>

Reviewed: June 2017

Published: March 3, 2020

Lister Hill National Center for Biomedical Communications
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National Institutes of Health
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