Opis choroby *

Definicja

The hereditary dentin disorders, dentinogenesis imperfecta (DGI) and dentin dysplasia (DD), comprise a group of conditions characterized by abnormal dentin structure affecting either the primary or both the primary and secondary dentitions.

Dane

Klasyfikacja
Kategoria

Kod ORPHA
167759

Kod OMIM
-

Kod ICD10
K00.5

Kod ICD11
-

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*Źródło

orphanet