Opis choroby *

Definicja

A rare genetic odontal or periodontal disorder that represents a group of developmental conditions affecting the structure and clinical appearance of the enamel of all or nearly all the teeth in a more or less equal manner, and which may be associated with morphologic or biochemical changes elsewhere in the body.

Dane

Klasyfikacja
Choroba

Kod ORPHA
88661

Kod OMIM
617217

Kod ICD10
K00.5

Kod ICD11
LA30.6

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*Źródło

orphanet