## 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