

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

*Źródło

orphanet