Opis choroby *

Definicja

A moderately severe form of osteogenesis imperfecta characterized by increased bone fragility and low bone mass that clinically manifests from infancy as susceptibility to bone fractures, short stature, mild to moderate scoliosis in most, gray-blue or white sclera, and dentinogenesis imperfecta.

Dane

Klasyfikacja
Podtyp kliniczny

Synonimy
OI type 4
OI typu 4

Kod ORPHA
216820

Kod OMIM
616507

Kod ICD10
Q78.0

Kod ICD11
LD24.K0

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*Źródło

orphanet