

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	Synonimy
Podtyp kliniczny	Ol type 4 Ol typu 4

Kod ORPHA	Kod OMIM	Kod ICD10
216820	616507	Q78.0

Kod ICD11
LD24.K0

*Źródło

orphanet