

Dermatoosteoliza typu Kirghiziana

Kod Orpha: 1657 Kod OMIM: 221810

Opis choroby *

Definicja

A rare genetic disease characterized by infantile onset of recurrent skin ulcerations, arthralgias, fever, peri-articular fistulous osteolysis, oligodontia, nail dystrophy, and keratitis. The disease takes a self-limiting course in childhood but results in severe cicatrization, chronic arthroses, pseudoacromegalic appearance of hands and feet, secondary scoliosis, and visual impairment. There have been no further descriptions in the literature since 1983.

Dane

Klasyfikacja

Zespół wad wrodzonych

Kod ORPHA

1657

Kod OMIM

221810

Kod ICD10

Q82.8

Kod ICD11

LD27.0Y

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.