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

Kod OMIM 221810

Kod ICD10 Q82.8

Kod ICD11 LD27.0Y

1657

<u>*Źródło</u>

orphanet