Opis choroby *

Definicja

An association reported in a single kindred characterized by the variable presence of the following features: anetodermia (macular atrophy of the skin), multiple exostoses, and brachydactyly type E. There have been no further descriptions in the literature since 1985.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
1962

Kod OMIM
133690

Kod ICD10
Q87.5

Kod ICD11
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*Źródło

orphanet