Opis choroby *

Definicja

A rare, genetic, skin tumor disorder characterized by childhood-onset of multiple, benign, asymptomatic, white to flesh-colored papules predominently located on the face, ears, neck and trunk, not associated with systemic organ involvement, associated malignancies or FLCN gene locus mutation.

Dane

Klasyfikacja
Choroba

Synonimy
Familial multiple trichodiscomas
Familial multiple trichodiscomas

Kod ORPHA
538756

Kod OMIM
190340

Kod ICD10
D23.9

Kod ICD11
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*Źródło

orphanet