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 <i>FLCN</i>gene locus mutation.

Dane

Klasyfikacja Synonimy

Choroba Familial multiple trichodiscomas

Familial multiple trichodiscomas

Kod ORPHA Kod OMIM Kod ICD10

538756 190340 D23.9

Kod ICD11

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*Źródło

orphanet