Opis choroby *

Definicja

A rare association syndrome, reported in several members of two families to date, characterized by arterial dissection, occurring at an early age and presenting with a range of manifestations depending on the vascular territory involved (ex. headache, dysphasia, hemiparesis), in association with cystic medial necrosis and multiple lentigines (brown and black in color and mainly affecting the skin of the trunk and extremities).

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
1682

Kod OMIM
600459

Kod ICD10
Q87.8

Kod ICD11
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*Źródło

orphanet