Opis choroby *

Definicja

A rare pervasive developmental disorder characterized by the presence of a unilateral angioma on the face and autistic developmental problems including language delay and atypical social interactions. The disease may initially resemble Sturge-Weber syndrome.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
137911

Kod OMIM
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Kod ICD10
F84.8

Kod ICD11
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*Źródło

orphanet