Opis choroby *

Definicja

A rare syndromic neurological disorder characterized by the association of Möbius syndrome (congenital facial palsy with impaired ocular abduction) with peripheral axonal neuropathy and hypogonadotropic hypogonadism. There have been no further reports since 1996.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
2560

Kod OMIM
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Kod ICD10
E23.0

Kod ICD11
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*Źródło

orphanet