Opis choroby *

Definicja

A syndrome characterized by unilateral or bilateral coronal synostosis, facial asymmetry, ptosis, strabismus and small ears with prominent superior and/or inferior crus, among other less common manifestations.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
ACS3
ACS 3
Akrocefalopolisyndaktylia typ 3
Acrocephalosyndactyly type 3
SCS

Kod ORPHA
794

Kod OMIM
101400

Kod ICD10
Q87.0

Kod ICD11
LD24.GY

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*Źródło

orphanet