Opis choroby *

Definicja

An acrocephalosyndactyly associated with craniosynostosis, midfacial hypoplasia, hand and foot malformation with a wide range of clinical expression and severity. Most of the affected patients show various other associated manifestations.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
ACS5
ACS 5
Akrocefalosyndaktylia typu 5
Acrocephalosyndactyly type 5

Kod ORPHA
710

Kod OMIM
101600

Kod ICD10
Q87.0

Kod ICD11
LD24.G0

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*Źródło

orphanet