

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 **Synonimy**

Zespół wad wrodzonych ACS5
 ACS 5
 Akrocefalo syndaktylia typu 5
 Acrocephalo syndactyly type 5

Kod ORPHA

710

Kod OMIM

101600

Kod ICD10

Q87.0

Kod ICD11

LD24.G0

*Źródło

orphanet