

## 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

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych ACS3	ACS 3
	Akrocefalopolisyndaktylia typ 3
	Acrocephalosyndactyly type 3
	SCS

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
794	101400	Q87.0

**Kod ICD11**  
LD24.GY

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### \*Źródło

orphanet