

## Opis choroby \*

### Definicja

A rare syndromic trigonocephaly characterized by marked malformations of the head and face (essentially acrocephaly), broad depressed nasal bridge, narrow maxillae, abnormalities of the hands and feet (polydactyly, brachydactyly, syndactyly, clinodactyly, camptodactyly, ulnar deviation), obesity and congenital heart disease. This disease is considered a variant of Carpenter syndrome without intellectual disability. There have been no further descriptions in the literature since 1992.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych ACPS4	ACPS4
	Akrocefalopolisyndaktylia typu 4
	Acrocephalopolysyndactyly type 4

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
65798	201020	Q87.0

### Kod ICD11

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### \*Źródło

orphanet