

## Opis choroby \*

### Definicja

A rare multisystemic genetic disorder characterized by a characteristic facial features with macrocephaly, overgrowth in infancy, intellectual disability and behavioral problems including anxieties and aggressiveness.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Sotos syndrome 2
	Zespół Sotosa 2

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
420179	614753	Q87.3

**Kod ICD11**  
LD2C

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### \*Źródło

orphanet