

## Opis choroby \*

### Definicja

Craniosynostosis-intracranial calcifications syndrome is a form of syndromic craniosynostosis characterized by pancraniosynostosis, head circumference below the mid-parental head circumference, mild facial dysmorphism (prominent supraorbital ridges, mild proptosis and maxillary hypoplasia) and calcification of the basal ganglia. The disease is associated with a favorable neurological outcome, normal intelligence and is inherited in an autosomal recessive manner.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Longman-Tolmie syndrome
	Zespół Longmana i Tolmiego

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
52054	608432	Q87.0

**Kod ICD11**

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### \*Źródło

orphanet