

## Opis choroby \*

### Definicja

A rare acrofacial dysostosis due to the presence of manifestations not usually seen in Nager syndrome (NS) such as microcephaly, blepharophimosis, microtia, a peculiar beaked nose, cleft lip and palate, symmetrical involvement of the thumbs and great toes and developmental delay. It has since been suggested that these features can also be a part of the NS phenotype.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Kennedy-Teebi syndrome Zespół Kennedy'ego i Teebiego

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
64542	-	Q75.4

**Kod ICD11**  
LD25.2

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### \*Źródło

orphanet