

## Opis choroby \*

### Definicja

Interstitial 9q22.3 microdeletion is associated with a phenotype including macrocephaly, overgrowth and trigonocephaly. Psychomotor delay, hyperactivity and distinctive facial features were also observed. It has been described in two unrelated children.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Microdeletion 9q22.3
	Mikrodelecja 9q22.3

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
77301	-	Q93.5

**Kod ICD11**  
LD44.90

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### \*Źródło

orphanet