

## Opis choroby \*

### Definicja

A drug-related embryofetopathy that can occur when an embryo/fetus is exposed to trimethadione and that is characterized by pre- and post-natal growth retardation, intellectual deficit, developmental and speech delay, craniofacial anomalies (with some similarities to those seen in fetal valproate syndrome), and less commonly, cleft palate, malformations of the heart, urogenital system and limbs. Trimethadione is an antiepileptic drug that has been removed from the market in Europe and is no longer used much in other countries due to teratogenicity and potential side effects.

### Dane

### Klasyfikacja

Zespół wad wrodzonych

#### Kod ORPHA

1913

#### Kod OMIM

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#### Kod ICD10

Q86.8

#### Kod ICD11

LD2F.0Y

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### \*Źródło

orphanet