

Embriopatia spowodowana talomidem

Kod Orpha: 3312 Kod OMIM:

Opis choroby *

Definicja

Thalidomide embryopathy is a group of anomalies presented in infants as a result of *in utero* exposure (between 20-36 days after fertilization) to thalidomide, a sedative used in treatment of a range of conditions, including morning sickness, leprosy and multiple myeloma (see these terms). Thalidomide embryopathy is characterized by phocomelia, amelia, forelimb and hand plate anomalies (absence of humerus and/or forearm, femur and/or lower leg, thumb anomalies). Other anomalies include facial hemangiomas, and damages to ears (anotia, microtia), eyes (microphthalmia, anophthalmos, coloboma, strabismus), internal organs (kidney, heart, and gastrointestinal tract), genitalia, and heart. Infant mortality associated with thalidomide embryopathy is estimated to be as high as 40%. Thalidomide is contraindicated in pregnancy and pregnancy prevention is recommended in women under treatment.

Dane

Klasyfikacja

Zespół wad wrodzonych

Synonimy

Fetal thalidomide syndrome
Płodowy zespół talomidowy

Kod ORPHA

3312

Kod OMIM

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Kod ICD10

Q86.8

Kod ICD11

LD2F.0Y

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

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