

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	Synonimy
Zespół wad wrodzonych	Fetal thalidomide syndrome
	Płodowy zespół talomidowy

Kod ORPHA	Kod OMIM	Kod ICD10
3312	-	Q86.8
Kod ICD11		
LD2F.0Y		

*Źródło

orphanet