

Systemowa przetoka tętniczo-żylna

Kod Orpha: 2039 Kod OMIM:

Opis choroby *

Definicja

Congenital systemic arteriovenous fistula is a rare, potentially life-threatening, vascular malformation characterized by a direct communication between an artery and a vein, without the interposition of the capillary bed, occurring in the systemic circulation (mainly the cranium, liver, lungs, extremities, and vessels in or near the thoracic wall). Manifestations are variable depending on size and extent of the fistula, the involved blood vessels and the precise location of the collaterals and may include systolic or continuous murmur over the affected organ, tachycardia, increased stroke volume, cardiomegaly and increased pulmonary vascular markings.

Dane

Klasyfikacja

Wada morfologiczna

Kod ORPHA
2039

Kod OMIM
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Kod ICD10
Q27.3

Kod ICD11
LA90.3Z

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

Dostępna na stronie www.orphanet.pl