

Opis choroby *

Definicja

A rare vascular anomaly characterized by the segmental narrowing of the abdominal and/or distal descending thoracic aorta, with varying involvement of the visceral and renal arteries, that commonly presents in children and young adults with early onset and refractory hypertension, abdominal angina, and lower-limb claudication, that can lead to life-threatening complications associated with severe hypertension (i.e. myocardial infarction, heart failure, aortic rupture, renal insufficiency and intracranial hemorrhage). It may be due to various congenital or acquired causes, but it is most often secondary to an acquired inflammatory disease (i.e. Takayasu arteritis or giant cell arteritis).

Dane

Klasyfikacja	Synonimy
Podtyp kliniczny	Coarctation of the abdominal aorta Koarktacja aorty brzusznej Zespół dysplastycznej aorty brzusznej Zespół zwężenia aorty brzusznej Mid-aortic dysplastic syndrome Mid-aortic syndrome Midaortic syndrome Middle aortic syndrome

Kod ORPHA	Kod OMIM	Kod ICD10
1456	-	Q25.1

Kod ICD11

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*Źródło

orphanet