Opis choroby *

Definicja

A number of families have been described, where several members were affected with coarctation of aorta. In a systematic study of coarctation, familial aggregation was considered as result of multifactorial inheritance and recurrence risks in sibs was evaluated at about 0.5% for coarctation and 1.0% for any form of congenital heart defect. Nevertheless, in some of the described families, aortic coarctations seems to be inherited as an autosomal dominant mutation.

Dane

Klasyfikacja
Podtyp kliniczny

Kod ORPHA
1455

Kod OMIM
-

Kod ICD10
Q25.1

Kod ICD11
-

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*Źródło

orphanet