

Autosomalnie dominująca koarktacja aorty

Kod Orpha: 1455 Kod OMIM:

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

-

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.