

## 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

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Kod ICD10

Q25.1

Kod ICD11

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### \*Źródło

orphanet