

Opis choroby *

Definicja

Kallmann syndrome with cardiopathy is characterised by hypogonadotropic hypogonadism associated with gonadotropin-releasing hormone (GnRH) deficiency, anosmia or hyposmia (with hypoplasia or aplasia of the olfactory bulbs) and complex congenital cardiac malformations (double-outlet right ventricle, dilated cardiomyopathy, right aortic arch). It represents a distinct clinical entity from Kallmann syndrome.

Dane

Klasyfikacja

Zespół wad wrodzonych

Kod ORPHA

2326

Kod OMIM

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Kod ICD10

Q24.8

Kod ICD11

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*Źródło

orphanet