

Opis choroby *

Definicja

A rare congenital non-syndromic heart malformation characterized by an abnormal protrusion of the interatrial septum into the right or left atrium, or both, during the cardiorespiratory cycle. The defect may be limited to the fossa ovalis or involve the entire septum. It can present as an isolated finding but is more often associated with interatrial shunts, in particular patent foramen ovale. Clinically it increases the risk of peripheral arterial embolism and stroke.

Dane

Klasyfikacja

Wada morfologiczna

Kod ORPHA	Kod OMIM	Kod ICD10
99107	-	Q21.1
Kod ICD11		
LA8E.Y		

*Źródło

orphanet