Opis choroby *

Definicja

Pulmonary arterial hypertension (PAH) is a group of diseases characterized by elevated pulmonary arterial resistance leading to right heart failure. PAH is progressive and potentially fatal. PAH may be idiopathic and/ or familial, or induced by drug or toxin (drug-or toxin-induced PAH, see these terms) or associated with other diseases like congenital heart disease, connective tissue disease, HIV, schistosomiasis, portal hypertension (PAH associated with other disease, see this term).

Dane

Klasyfikacja
Kategoria

Synonimy
PAH
PAH

Kod ORPHA
182090

Kod OMIM
-

Kod ICD10
-

Kod ICD11
BB01.0

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*Źródło

orphanet