Opis choroby *

Definicja

A rare hematologic disease characterized by eosinophilia without evidence of clonality persisting for at least six months, for which no underlying cause can be identified. The condition is associated with signs of organ damage and dysfunction. Clinical manifestations are highly variable, depending on the organ systems involved, and include rapidly developing, life-threatening cardiovascular or neurological complications.

Dane

Klasyfikacja
Choroba

Kod ORPHA
3260

Kod OMIM
607685

Kod ICD10
D47.5

Kod ICD11
4B03.Z

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*Źródło

orphanet