

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

*Źródło

orphanet