Opis choroby *

Definicja

A rare hypereosinophilic syndrome characterized by hypereosinophilia produced by clonal eosinophils derived from neoplastic stem cells in the absence of any secondary cause of eosinophilia and persisting for at least six months. The condition is associated with signs of organ infiltration, dysfunction, and damage. Clinical manifestations are highly variable, depending on the organ systems involved, and include dermatologic, pulmonary, cardiac, gastrointestinal, and cerebral manifestations, among others.

Dane

Klasyfikacja
Choroba

Synonimy
Clonal hypereosinophilic syndrome
HES-M
HES-N
Klonalny zespół hipereozynofilowy
Neoplastyczny zespół hipereozynofilowy
Pierwotny HES
HES-M
HES-N
Neoplastic hypereosinophilic syndrome
Primary HES

Kod ORPHA
314950

Kod OMIM
-

Kod ICD10
D47.5

Kod ICD11
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*Źródło

orphanet