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 Synonimy

Choroba 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 Kod OMIM Kod ICD10

314950 - D47.5

Kod ICD11

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*Źródło

orphanet