Opis choroby *

Definicja

A rare subtype of indolent systemic mastocytosis characterized by isolated bone marrow involvement without skin lesions, low burden of neoplastic mast cells, and often normal or near normal serum tryptase levels. The KIT D816V mutation is present in the majority of cases.

Dane

Klasyfikacja
Choroba

Kod ORPHA
158778

Kod OMIM
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Kod ICD10
D47.0

Kod ICD11
2A21.0Y

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*Źródło

orphanet