

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

*Źródło

orphanet