Opis choroby *

Definicja

A rare group of mastocytosis diseases characterized by abnormal accumulation and proliferation of mast cells in the skin and including the three recognised forms: diffuse cutaneous mastocytosis, cutaneous mastocytoma and, the most common form, maculopapular cutaneous mastocytosis. In some cases (most commonly in adults), cutaneous mastocytosis may occur in association with mast cell infiltration of various extracutaneous organs, in which case the disorder is referred to as systemic mastocytosis.

Dane

Klasyfikacja
Grupa fenomenów

Kod ORPHA
66646

Kod OMIM
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Kod ICD10
Q82.2

Kod ICD11
2A21.1

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*Źródło

orphanet