Opis choroby *

Definicja

A rare, systemic disease with skin involvement characterized by the onset of idiopathic lupus erythematosus-like signs and symptoms resulting from continuous drug intake (>1 month), which resolve when treatment is discontinued, in persons with no history of autoimmune disease. Manifestations are variable and may be systemic (e.g. arthralgia, myalgia, fever, fatigue, serositis, pleuritis, pericarditis), subacute cutaneous (incl. photosensitive, non-scarring, annular, polycyclic or papulosquamous lesions, malar erythema, vasculitis, bullous lesions, erythema multiforme-like changes), and/or chronic cutaneous (typically discoid lesions in sun-exposed areas). Procainamide and hydralazine are the drugs most frequently implicated.

Dane

Klasyfikacja
Choroba

Synonimy
DILE
DILE

Kod ORPHA
231111

Kod OMIM
-

Kod ICD10
M32.0

Kod ICD11
4A40.1

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*Źródło

orphanet