Opis choroby *

Definicja

A rare systemic autoimmune disease characterized by the presence of signs and symptoms suggestive of a systemic autoimmune disease that do not fulfil the existing classification criteria. The main clinical manifestations are arthritis with arthralgia, Raynaud's phenomenon, xerostomia, xerophthalmia, and leukopenia, while neurologic or renal involvement are virtually absent.

Dane

Klasyfikacja
Choroba

Synonimy
UCTD
UCTD

Kod ORPHA
90002

Kod OMIM
-

Kod ICD10
M35.8

Kod ICD11
LD28.Y

---

*Źródło

orphanet