Opis choroby *

Definicja

A rare idiopathic interstitial pneumonia characterized by temporally uniform alveolar and interstitial mononuclear cell inflammation (cellular type) and/or fibrosis of the alveolar walls (fibrotic type) with preserved alveolar architecture. Other types of interstitial lung disease must be excluded. Symptoms are non-specific and include dyspnea, cough, and often constitutional symptoms such as fever and fatigue. Pulmonary function test reveals a restrictive pattern. Computed tomography shows predominantly lower lobe subpleural reticular changes, traction bronchiectasis, and ground-glass opacities. The cellular type of the disease is less common but carries a better prognosis.

Dane

Klasyfikacja
Choroba

Synonimy
NSIP
Niespecyficzne idiopatyczne śródmiąższowe zapalenie płuc
NSIP
Non-specific idiopathic interstitial pneumonia

Kod ORPHA
91364

Kod OMIM
-

Kod ICD10
J84.8

Kod ICD11
CB03.Y

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*Źródło

orphanet