Opis choroby *

Definicja

A rare, genetic, interstitial lung disease due to mutations in the CSF2R (colony-stimulating factor 2 receptor) alpha or beta subunits and characterized by alveolar accumulation of pulmonary surfactant, presenting a highly variable clinical presentation, ranging from asymptomatic to severe respiratory failure. Characteristic lung biopsy findings include periodic acid-Schiff-positive, granular eosinophilic material, enlarged foamy alveolar macrophages, and well-preserved alveolar walls. The Granulocyte-macrophage colony-stimulating factor (GM-CSF) receptor function is impaired but GM-CSF receptor autoantibodies are absent.

Dane

Klasyfikacja
Choroba

Synonimy
Congenital PAP
Wrodzona PAP
Congenital pulmonary alveolar proteinosis

Kod ORPHA
264675

Kod OMIM
300770

Kod ICD10
J84.0

Kod ICD11
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*Źródło

orphanet