## Opis choroby *

Definicja

A rare primary interstitial lung disease characterized by the accumulation of lipids and proteins related to surfactant in the alveoli in association with the presence of antibodies against granulocyte-macrophage colony-stimulating factor (GM-CSF). The disease leads to a progressive impairment of gas exchange and respiratory insufficiency.

Dane

**Klasyfikacja**
Choroba

Synonimy
Autoimmune PAP
aPAP
Autoimmunologiczna PAP
Idiopatyczna PAP
Idiopatyczna proteinoza pęcherzyków płucnych
iPAP
aPAP

**Kod ORPHA**
747

**Kod OMIM**
610910

**Kod ICD10**
J84.0

**Kod ICD11**
CB04.31

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*Źródło

orphanet