Opis choroby *

Definicja

A rare, acquired, interstitial lung disease, characterized by alveolar surfactant accumulation, cough, progressive dyspnea and respiratory insufficiency. The disease may be secondary to hematological disorder, toxic inhalation, and infection or may occur within the setting of immunosuppression after transplantation.

Dane

Klasyfikacja
Choroba

Synonimy
Secondary PAP
Wtórna PAP

Kod ORPHA
420259

Kod OMIM
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Kod ICD10
J84.0

Kod ICD11
CB04.31

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*Źródło

orphanet