Opis choroby *

Definicja

A group of interstitial lung diseases (ILD) induced by genetic mutations disrupting surfactant function and gas exchange in the lung. The disorders caused by these mutations affect full-term infants and older children and exhibit considerable overlap in their clinical and histologic presentation.

Dane

Klasyfikacja
Kategoria

Synonimy
Primary ILD specific to childhood due to pulmonary surfactant protein anomalies
Pierwotna dziecięca ILD spowodowana anomaliami białek surfaktantu

Kod ORPHA
100049

Kod OMIM
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Kod ICD10
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Kod ICD11
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*Źródło

orphanet