

## 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 anomaliasi 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