## Opis choroby \*

## Definicja

A rare genetic systemic or rheumatologic disease characterized by interstitial lung disease (often with pulmonary hemorrhage) and inflammatory arthritis, associated with high-titer autoantibodies (including anti-nuclear and anti-neutrophil cytoplasmic antibodies, and rheumatoid factor). Patients present from infancy to adolescence with tachypnea, cough, hemoptysis, and/or joint pain. Some patients may also develop glomerular disease.

Dane

**Klasyfikacja** Synonimy

Choroba COPA syndrome COPA syndrome

 Kod ORPHA
 Kod OMIM
 Kod ICD10

 444092
 616414
 J84.8

**Kod ICD11** 

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## \*Źródło

orphanet