# DISEASE:
# Autoimmune interstitial lung disease-arthritis syndrome

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| **NAME:** | Autoimmune interstitial lung disease-arthritis syndrome |
| **DESCRIPTION:** | 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. |
| **ORPHACODE:** | 444092 |
| **SYNONYMS:** | COPA syndrome |
| **XREF(S):** | Orphanet<br>ICD-10<br>OMIM |
| **ANALYTE(S):** | COPA |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

# RELATED CONTENT

## Related Genetic Tests

- Child Interstitial Lung Disease (child - gene panel)

## Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

## Related Analytes

- COPI coat complex subunit alpha

## Related Gene Panels

- chILD (34 genes) - KUL