

## Opis choroby \*

### Definicja

A rare renal disease occurring in the setting of a systemic IgG4 related disease (IgG4-RD). The disorder is characterized by a fibrosing tubulointerstitial nephritis consisting of predominantly IgG4+ plasma cells with/without glomerulonephritis, retroperitoneal fibrosis and hydronephrosis.

### Dane

### Klasyfikacja

Podtyp kliniczny

#### Kod ORPHA

449395

#### Kod OMIM

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#### Kod ICD10

N11.8

#### Kod ICD11

4A43.0

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### \*Źródło

orphanet