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
-

Kod ICD10
N11.8

Kod ICD11
4A43.0

---

*Źródło

orphanet