Opis choroby *

Definicja

A rare monoclonalgammopathy characterized by renal proximal tubule dysfunction secondary to monoclonal kappa light chain deposits in proximal tubular cells. Clinical presentation is with variable chronic kidney disease, low molecular weight proteinuria, aminoaciduria, hyperphosphaturia, uricosuria, bicarbonaturia, and non-diabetic glycosuria. Renal phosphate and urate wasting may cause hypophosphatemia and hypouricaemia.

Dane

Klasyfikacja
Choroba

Synonimy
Acquired Fanconi syndrome secondary to monoclonal gammopathy
Zespół Fanconiego ze szpiczakiem
Zespół Fanconiego związany z lekkimi łańcuchami kappa
Acquired monoclonal immunoglobulin light chain-associated Fanconi syndrome

Kod ORPHA
91136

Kod OMIM
-

Kod ICD10
E72.0

Kod ICD11
-

---

*Źródło

orphanet