

## 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

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#### Kod ICD10

E72.0

#### Kod ICD11

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#### [\\*Źródło](#)

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