

Rodzinna glikozuria nerkowa

Kod Orpha: 69076 Kod OMIM: 233100

Opis choroby *

Definicja

A rare, genetic, glucose transport disorder characterized by the presence of persistent isolated glucosuria in the absence of both proximal tubular dysfunction and hyperglycemia. The disorder is benign in the majority of cases although it may occasionally manifest with polyuria, enuresis, a mild growth and pubertal maturation delay, hypercalciuria, aminoaciduria and, in severe cases, increased incidence of urinary infections and episodic dehydration and ketosis during pregnancy and starvation.

Dane

Klasyfikacja

Choroba

Synonimy

Familial renal glycosuria
Niedobór SGLT2
SGLT2 deficiency

Kod ORPHA

69076

Kod OMIM

233100

Kod ICD10

E74.8

Kod ICD11

5C61.5

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.