

Rodzinny hipoaldosteronizm

Kod Orpha: 427 Kod OMIM: 610600

Opis choroby *

Definicja

A rare genetic hypoaldosteronism that typically presents in infancy (early-onset familial hypoaldosteronism) as a life-threatening electrolyte imbalance (failure to thrive, recurrent vomiting, and severe dehydration). A history of fever, diarrhoea, lethargy, poor weight gain, poor feeding since birth may also be present. Older subjects (late-onset familial hypoaldosteronism) are less severely affected or asymptomatic.

Dane

Klasyfikacja

Choroba

Kod ORPHA

427

Kod OMIM

610600

Kod ICD10

E27.4

Kod ICD11

5A73

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.