

## 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

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### \*Źródło

orphanet